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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<p>(21) International Application Number: PCT/US96/20744</p> <p>(22) International Filing Date: 20 December 1996 (20.12.96)</p> <p>(30) Priority Data: 60/009,532 29 December 1995 (29.12.95) US</p> <p>(71) Applicant (for all designated States except US): SMITHKLINE BEECHAM CORPORATION [US/US]; Corporate Intellectual Property, UW2220, 709 Swedeland Road, P.O. Box 1539, King of Prussia, PA 19406-0939 (US).</p> <p>(72) Inventors; and</p> <p>(75) Inventors/Applicants (for US only): ALI, Fadia, E. [US/US]; 5 North Green Acre Drive, Cherry Hill, NJ 08003 (US). BONDINELL, William, E. [US/US]; 1512 Franklin Lane, Wayne, PA 19087 (US). KEENAN, Richard, M. [US/US]; 796 Bass Cove, Malvern, PA 19355 (US). KU, Thomas, Wen, Fu [US/US]; 1413 Southwind Way, Dresher, PA 19025 (US). MILLER, William, H. [US/US]; 333 Fell Lane, Schwenksville, PA 19473 (US). SAMANEN, James [US/US]; 145 Jug Hollow Road, Phoenixville, PA 19460 (US).</p>		<p>(74) Agents: KINZIG, Charles, M. et al.; SmithKline Beecham Corporation, Corporate Intellectual Property, UW2220, 709 Swedeland Road, P.O. Box 1539, King of Prussia, PA 19406-0939 (US).</p> <p>(81) Designated States: AL, AM, AU, BB, BG, BR, CA, CN, CZ, EE, GE, HU, IL, IS, JP, KG, KP, KR, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, SG, SI, SK, TR, TT, UA, US, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p>Published <i>With international search report.</i></p>
<p>(54) Title: VITRONECTIN RECEPTOR ANTAGONISTS</p> <p>(57) Abstract</p> <p>Compounds of formula (I) are disclosed which are vitronectin receptor antagonists useful in the treatment of osteoporosis.</p> <div style="text-align: center;"> <p style="text-align: right;">(I)</p> </div>		

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TITLE
Vitronectin Receptor Antagonists

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FIELD OF THE INVENTION

This invention relates to pharmaceutically active compounds which inhibit the vitronectin receptor and are useful for the treatment of diseases wherein inhibition of the vitronectin receptor is indicated, such as inflammation, cancer, angiogenesis, atherosclerosis, restenosis, and diseases wherein bone resorption is a factor.

10

BACKGROUND OF THE INVENTION

Integrins are a superfamily of cell adhesion receptors, which are transmembrane glycoproteins expressed on a variety of cells. These cell surface adhesion receptors include gpIIb /IIIa, the fibrinogen receptor, and $\alpha_v\beta_3$, the vitronectin receptor. The fibrinogen receptor gpIIb /IIIa is expressed on the platelet surface and it mediates platelet aggregation and the formation of a hemostatic clot at the site of a bleeding wound. Philips, *et al.*, *Blood.*, **1988**, *71*, 831.

The vitronectin receptor $\alpha_v\beta_3$ is expressed on a number of cells, including endothelial, smooth muscle, osteoclast, and tumor cells, and, thus, it has a variety of functions. The $\alpha_v\beta_3$ receptor expressed on the membrane of osteoclast cells is believed to play a role in the bone resorption process and contribute to the development of osteoporosis. Ross, *et al.*, *J. Biol. Chem.*, **1987**, *262*, 7703; Fisher, *et al.*, *Endocrinology* **1993**, *132*, 1411; Bertolini *et al.*, *J. Bone Min. Res.*, *6*, Sup. 1, S146, 252; EP 528 587 and 528 586. The $\alpha_v\beta_3$ receptor expressed on human aortic smooth muscle cells stimulates their migration into neointima, which leads to the formation of atherosclerosis and restenosis after angioplasty. Brown, *et al.*, *Cardiovascular Res.*, **1994**, *28*, 1815. Additionally, a recent study has shown that a $\alpha_v\beta_3$ antagonist is able to promote tumor regression by inducing apoptosis of angiogenic blood vessels. Brooks, *et al.*, *Cell*, **1994**, *79*, 1157. Thus, agents that would block the vitronectin receptor would be useful in treating diseases mediated by this receptor, such as osteoporosis, atherosclerosis, restenosis and cancer.

Alig et al., EP 0 381 033, Hartman, et al., EP 0 540,334, Blackburn, et al., WO 93/08174, Bondinell, et al., WO 95/18619, Bondinell, et al., WO 94/14776, Blackburn, et al. WO 95/04057, Egbertson, et al, EP 0 478 328, Sugihara, et al. EP 529,858, Porter, et al., EP 0 542 363, and Fisher, et al., EP 0 635 492, and many others disclose certain
5 compounds that are useful for selectively inhibiting the fibrinogen receptor. PCT/US95/08306, filed June 29, 1995 (SmithKline Beecham Corp.) and PCT/US95/08146 filed June 29, 1995 (SmithKline Beecham Corp.) disclose vitronectin receptor selective antagonists. However, there are few reports of compounds which are potent vitronectin receptor antagonists. It has now been discovered that certain
10 appropriately substituted amino pyridine compounds are potent inhibitors of the vitronectin receptor. In particular, it has been discovered that the amino pyridine moiety may be combined with a fibrinogen antagonist template to prepare compounds which are more potent inhibitors of the vitronectin receptor than the fibrinogen receptor.

15 SUMMARY OF THE INVENTION

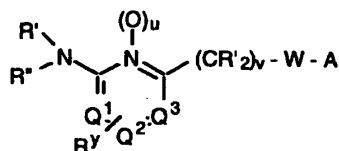
This invention comprises compounds of the formula (I) as described hereinafter, which have pharmacological activity for the inhibition of the vitronectin receptor and are useful in the treatment of inflammation, cancer, cardiovascular disorders, such as atherosclerosis and restenosis, and diseases wherein bone resorption is a factor, such as
20 osteoporosis.

This invention is also a pharmaceutical composition comprising a compound according to formula (I) and a pharmaceutically acceptable carrier.

This invention is also a method of treating diseases which are mediated by the vitronectin receptor. In a particular aspect, the compounds of this invention are useful for
25 treating atherosclerosis, restenosis, inflammation, cancer and osteoporosis.

DETAILED DESCRIPTION

This invention comprises novel compounds which are more potent inhibitors of the vitronectin receptor than the fibrinogen receptor. The compounds of the instant
30 invention comprise a fibrinogen receptor antagonist template that is linked to an optionally substituted *o*-amino pyridine moiety according to formula (I):



(I)

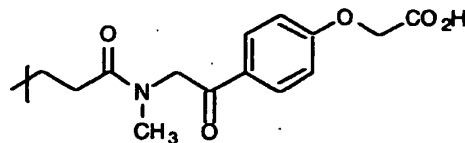
wherein

- 5 A is a fibrinogen antagonist template;
W is a linking moiety of the form $-(\text{CHR}^g)_a-\text{U}-(\text{CHR}^g)_b-\text{V}-$;
Q¹, Q² and Q³ are independently N or C-R^y, provided that no more than one of Q¹, Q²
and Q³ is N;
R' is H or C₁₋₆alkyl, C₃₋₇cycloalkyl-C₀₋₆alkyl or Ar-C₀₋₆alkyl
10 R'' is R', -C(O)R' or -C(O)OR';
R^g is H or C₁₋₆alkyl, Het-C₀₋₆alkyl, C₃₋₇cycloalkyl-C₀₋₆alkyl or Ar-C₀₋₆alkyl;
R^k is R^g, -C(O)R^g or -C(O)OR^g
Rⁱ is H, C₁₋₆alkyl, Het-C₀₋₆alkyl, C₃₋₇cycloalkyl-C₀₋₆alkyl, Ar-C₀₋₆alkyl, Het-C₀₋₆alkyl-
U'-C₁₋₆alkyl, C₃₋₇cycloalkyl-C₀₋₆alkyl-U'-C₁₋₆alkyl or Ar-C₀₋₆alkyl-U'-C₁₋₆alkyl;
15 R^y is H, halo, -OR^g, -SR^g, -CN, -NR^gR^k, -NO₂, -CF₃, CF₃S(O)_r-, -CO₂R^g, -COR^g or
-CONR₂;
U and V are absent or CO, CR₂, C(=CR₂), S(O)_c, O, NR^g, CR^gOR^g, CR^g(OR^k)CR₂,
CR₂CR^g(OR^k), C(O)CR₂, CR₂C(O), CONRⁱ, NRⁱCO, OC(O), C(O)O, C(S)O,
OC(S), C(S)NR^g, NR^gC(S), S(O)₂NR^g, NR^gS(O)₂, N=N, NR^gNR^g, NR^gCR₂,
20 NR^gCR₂, CR₂O, OCR₂, CR^g=CR^g, C≡C, Ar or Het;
a is 0, 1 or 2;
b is 0, 1 or 2;
c is 0, 1 or 2;
u is 0 or 1;
25 v is 0 or 1;
and pharmaceutically acceptable salts thereof;
provided that:
(i) when v is 0, and R', R'' and R' are H, and Q¹-Q³ are CH,
W-A is not 7-aminocarbonyl-2,3,4,5-tetrahydro-3-oxo-4-methyl-1H-1,4-
30 benzodiazepine-2-acetic acid, 7-aminocarbonyl-1-acetyl-2,3,4,5-tetrahydro-3-oxo-
4-methyl-1H-1,4-benzodiazepine-2-acetic acid, or 7-aminocarbonyl-2,3,4,5-

tetrahydro-3-oxo-4-methyl-1H-1-benzazepine-2-acetic acid, or the methyl esters thereof;

(ii) when v is 0 or 1 and R' , R'' and R' are H, and Q^1 - Q^3 are CH,

W-A is not 3-propanoyl-glycyl-aspartyl-phenylalanine, or



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, and the benzyl esters thereof.

Preferably, Q^1 , Q^2 and Q^3 are all CH, and u is 0.

Suitably, R' is H and R'' is H or C_{1-4} alkyl.

Preferably v is 1.

Suitably, W is $-(CHR^8)_a-CONR^i-$ or $-(CHR^8)_a-NR^iCO-$

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Suitably U' is $CONR'$ or $NR'CO$.

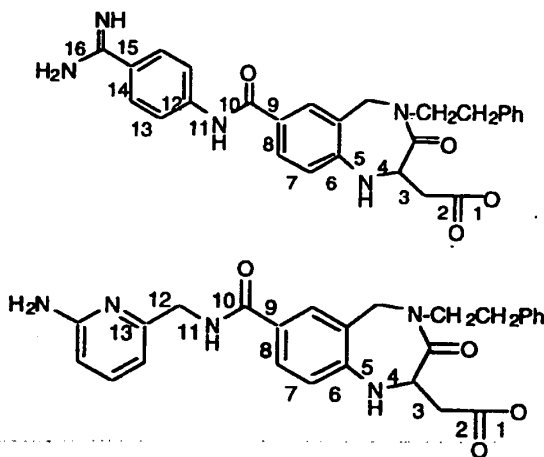
A fibrinogen receptor antagonist is an agent that inhibits the binding of fibrinogen to the platelet-bound fibrinogen receptor GPIIb-IIIa. Many fibrinogen antagonists are known to the art. As used herein, the term "fibrinogen receptor antagonist template" means the core structure of a fibrinogen receptor antagonist, said core containing an acidic group and being linked to an organic group substituted with a basic nitrogen moiety. Typically, the core structure adds some form of rigid spacing between the acidic moiety and the basic nitrogen moiety, and contains one or more ring structures or amide bonds to effect this. It is preferred that about twelve to fifteen, more preferably thirteen or fourteen, intervening covalent bonds via the shortest intramolecular path will exist between the acidic group of the fibrinogen receptor antagonist template and nitrogen of the pyridine moiety in formula (I). It is an object of this invention that a fibrinogen receptor antagonist is converted to a vitronectin receptor antagonist by replacing the basic nitrogen group in a fibrinogen receptor antagonist with an optionally substituted *o*-amino pyridine group. In addition, the number of intervening covalent bonds between the acidic moiety and the nitrogen of the pyridine will be about two to five, preferably three or four, covalent bonds shorter than the number of intervening covalent bonds between the acidic moiety and the basic nitrogen group of the fibrinogen antagonist. The identity of the linking moiety W may be chosen to obtain the proper spacing between the acidic moiety of the fibrinogen antagonist template and the nitrogen atom of the pyridine. Generally, a

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fibrinogen antagonist will have an intramolecular distance of about 16 angstroms between the acidic moiety (e.g., the atom which gives up the proton or accepts the electron pair) and the basic moiety (e.g., which accepts a proton or donates and electron pair), while the vitronectin antagonist will have about 14 angstroms between the respective acidic and basic centers.

For purposes of illustration, using the 7-2,3,4,5-tetrahydro-3-oxo-4-methyl-benzodiazepine fibrinogen antagonist template disclosed in WO 93/08174 as a suitable fibrinogen antagonist template, the compound (R,S)-7-[[[4-(aminoiminomethyl)phenyl]amino]carbonyl]-4-(2-phenylethyl)-1,3,4,5-tetrahydro-3-oxo-2H-1,4-benzodiazepine-2-acetic acid, which is potent and selective fibrinogen antagonist, is converted to a potent and selective vitronectin receptor antagonist by replacing the 4-(aminoiminomethyl)phenyl moiety with the 6-amino-pyrid-2-yl moiety. As illustrated below in Figure 1, in the former case, there are sixteen intervening covalent bonds between the acidic moiety and the basic moiety; in the fibrinogen antagonist whereas, in the latter case there are 13 intervening covalent bonds in the vitronectin antagonist of this invention.

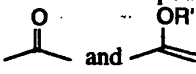
Figure 1



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In fact the 4-(aminoiminomethyl)phenyl moiety is a common substituent on fibrinogen antagonist templates known to the art, and simple replacement of this moiety with an optionally substituted (6-aminopyrid-2-yl)methyl moiety may serve as guide to

converting compounds having known fibrinogen antagonist templates into vitronectin receptor antagonists.

Also included in this invention are pharmaceutically acceptable addition salts, complexes or prodrugs of the compounds of this invention. Prodrugs are considered to be any covalently bonded carriers which release the active parent drug according to formula (I) *in vivo*. In cases wherein the compounds of this invention may have one or more chiral centers, unless specified, this invention includes each unique nonracemic compound which may be synthesized and resolved by conventional techniques. In cases in which compounds have unsaturated carbon-carbon double bonds, both the *cis* (Z) and *trans* (E) isomers are within the scope of this invention. In cases wherein compounds may exist in tautomeric forms, such as keto-enol tautomers, such as , and each tautomeric form is contemplated as being included within this invention whether existing in equilibrium or locked in one form by appropriate substitution with R'.

The compounds of formula (I) inhibit the binding of vitronectin and other RGD-containing peptides to the vitronectin ($\alpha_v\beta_3$) receptor. Inhibition of the vitronectin receptor on osteoclasts inhibits osteoclastic bone resorption and is useful in the treatment of diseases wherein bone resorption is associated with pathology, such as osteoporosis. Additionally, since the compounds of the instant invention inhibit vitronectin receptors on a number of different types of cells, said compounds would be useful in the treatment of inflammation and cardiovascular diseases, such as atherosclerosis and restenosis, and would be useful as anti-metastatic and antitumor agents.

Table I, below, describes certain fibrinogen receptor antagonists, whose core structures are useful in carrying out the instant invention. Reference should be made to the patent applications and other publications for their full disclosures, including the methods of preparing said templates and specific compounds embodying said templates. The entire disclosure of the noted patent applications and other publications is incorporated herein by reference as though fully set forth. The list following is not intended to limit the scope of the present invention, but only to illustrate certain known templates:

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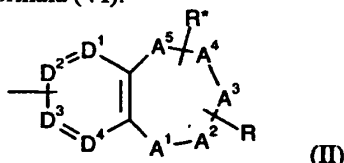
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In one particular embodiment, the fibrinogen receptor antagonist template A is the fused 6/7 ring bicyclic ring defined in Bondinell, et al., WO 93/00095, published January 7, 1993, as defined by sub-formula (VI):



15 wherein

A¹ to A⁵ form an accessible substituted seven-membered ring, which may be saturated or unsaturated, optionally containing up to two heteroatoms chosen from the group of O, S and N wherein S and N may be optionally oxidized;

20 D¹ to D⁴ form an accessible substituted six membered ring, optionally containing up to two nitrogen atoms;

R is at least one substituent chosen from the group of R⁷, or Q-C₁₋₄alkyl, Q-C₂₋₄alkenyl, Q-C₂₋₄alkynyl, optionally substituted by one or more of =O, R¹¹ or R⁷;

25 R* is H, Q-C₁₋₆alkyl, Q-C₁₋₆oxoalkyl, Q-C₂₋₆alkenyl, Q-C₃₋₄oxoalkenyl, Q-C₃₋₄oxoalkynyl, Q-C₂₋₄alkynyl, C₃₋₆cycloalkyl, Ar or Het, optionally substituted by one or more of R¹¹;

Q is H, C₃₋₆cycloalkyl, Het or Ar;

R⁷ is -COR⁸, -COCR'₂R⁹, -C(S)R⁸, -S(O)_mOR', -S(O)_mNR'R'', -PO(OR'), -PO(OR')₂, -B(OR')₂, -NO₂ and Tet;

30 R⁸ is -OR', -NR'R'', -NR'SO₂R', -NR'OR', -OCR'₂C(O)OR', -OCR'₂OC(O)-R', -OCR'₂C(O)NR'₂, CF₃ or AA¹;

R⁹ is -OR', -CN, -S(O)_rR', S(O)_mNR'₂, -C(O)R' C(O)NR'₂ or -CO₂R';

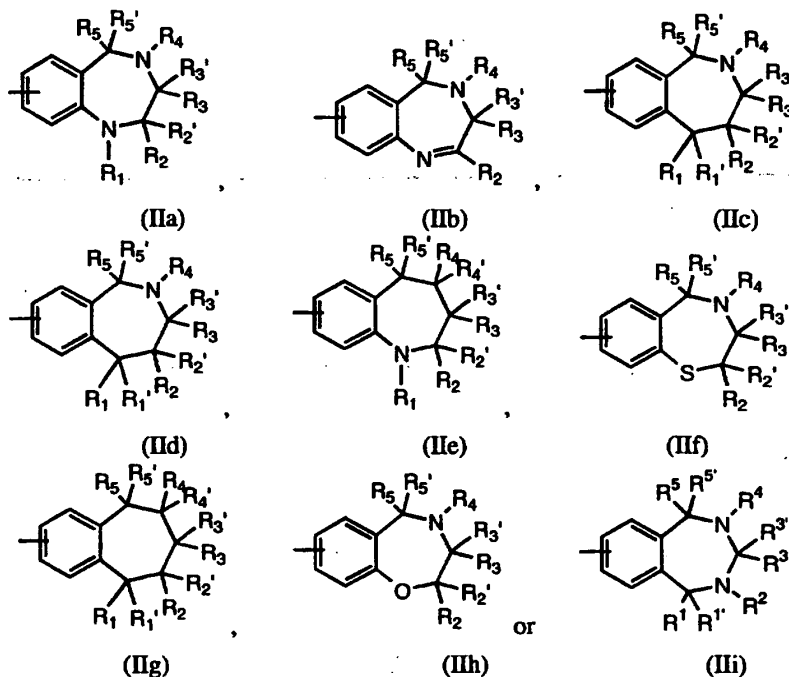
R¹¹ is H, halo, -OR¹², -CN, -NR'R¹², -NO₂, -CF₃, CF₃S(O)_r, -CO₂R', -CONR'₂, Q-C₀₋₆alkyl-, Q-C₁₋₆oxoalkyl-, Q-C₂₋₆alkenyl-, Q-C₂₋₆alkynyl-, Q-C₀₋₆alkyloxy-, Q-C₀₋₆alkylamino- or Q-C₀₋₆alkyl-S(O)_r;

- R^{12} is R' , $-C(O)R'$, $-C(O)NR'^2$, $-C(O)OR^{15}$, $-S(O)_mR'$ or $S(O)_mNR'^2$;
 R^{13} is R' , $-CF_3$, $-SR'$, or $-OR'$;
 R^{14} is R' , $C(O)R'$, CN , NO_2 , SO_2R' or $C(O)OR^{15}$;
 R^{15} is H , C_{1-6} alkyl or $Ar-C_{0-4}$ alkyl;
5 R' is H , C_{1-6} alkyl, C_{3-7} cycloalkyl- C_{0-4} alkyl or $Ar-C_{0-4}$ alkyl;
 R'' is R' , $-C(O)R'$ or $-C(O)OR^{15}$;
 R''' is R'' or $AA2$;
 $AA1$ is an amino acid attached through its amino group and having its carboxyl group optionally protected, and $AA2$ is an amino acid attached through its carboxyl group, and having its amino group optionally protected;
10 m is 1 or 2;
 n is 0 to 3;
 p is 0 or 1; and
 t is 0 to 2; or
15 pharmaceutically acceptable salts thereof.
With reference to formula (II), suitably,
 A^1 is $CR^1R^{1'}$, CR^1 , NR^1 , N , O or $S(O)_x$;
 A^2 is $CR^2R^{2'}$, CR^2 , NR^2 ;
 A^3 is $CR^3R^{3'}$, CR^3 , NR^3 , N , O or $S(O)_x$;
20 A^4 is $CR^4R^{4'}$, CR^4 , NR^4 , or N ;
 A^5 is $CR^5R^{5'}$, CR^5 , NR^5 , N , O or $S(O)_x$;
 D^1 - D^4 are CR^{11} , CR^6 or N ;
 R^1 and $R^{1'}$ are R^* or R , or together are $=O$;
 R^2 and $R^{2'}$ are R^* , R or $=O$;
25 R^3 and $R^{3'}$ are R^* , R or $=O$;
 R^4 and $R^{4'}$ are R^* , R or $=O$;
 R^5 and $R^{5'}$ are R^* , R or $=O$; and
 x is 0 to 2.
More suitably, A^1 is $CR^1R^{1'}$, CR^1 , NR^1 , N , O or S ; A^2 is $CR^2R^{2'}$, NR^2 or CR^2 ;
30 A^3 is $CR^3R^{3'}$; A^4 is $CR^4R^{4'}$, CR^4 , NR^4 , or N ; A^5 is $CR^5R^{5'}$, CR^5 , NR^5 , N , O ; D^1 - D^4 are CH ; R^2 or R^4 are R ; R^3 , $R^{3'}$ and R^5 , $R^{5'}$ are $=O$ or R^* , H .
Preferably, A^1 is CHR^1 , CR^1 , NR^1 , N or S ; A^2 is CR^2 or $CR^2R^{2'}$; A^3 is $CR^3R^{3'}$;
 A^4 is $CR^4R^{4'}$ or NR^4 ; A^5 is $CR^5R^{5'}$, and D^1 - D^4 are CH .
In one embodiment, A^1 is CR^1 , A^2 is CR^2 , A^3 is $C=O$, A^4 is NR^4 and A^5 are
35 CHR^5 .
In another embodiment, A^1 is NR^1 , A^2 is $CHCR^2$, A^3 is $CR^3R^{3'}$, A^4 is NR^4 , and A^5 are $C=O$.

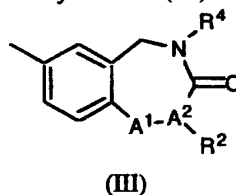
In yet another embodiment, A^1 and A^4 are $C=O$, A^2 is NR^2 , A^3 is CHR^3 and A^5 is NR^5 .

In a preferred embodiment, A^1 is NR^1 , A^2 is CHR^2 , A^3 is $C=O$, A^4 is NR' and A^5 is CHR^5 .

5 Representative sub-formulas of (II) are given by each of formulas (IIa)-(Iii) below:



A preferred template is given by formula (III):



wherein

A^1-A^2 is NR^1-CH , $NC(O)R^3-CH$, $N=C$, $CR^1=C$, CHR^1-CH , $O-CH$ or $S-CH$;

R^1 is H, C_{1-6} alkyl or benzyl;

20 R^2 is $(CH_2)_qCO_2H$;

R^4 is H, C_{1-6} alkyl, $Ar-C_{0-6}$ alkyl, $Het-C_{0-6}$ alkyl, or C_{3-6} cycloalkyl- C_{0-6} alkyl; and

q is 1, 2 or 3.

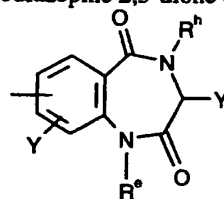
Preferably A¹-A² is NH-CH and R² is CH₂CO₂H. Suitably, R³ is methyl and W (as defined in formula (I)) is (CH₂)_aNR¹CO. Suitably R¹ is substituted by NHR', CN, CO₂H, biotin, benzimidazole or optionally substituted phenyl.

Specific examples of vitronectin antagonists employing this template are:

- 5 (S)-7-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic;
 (S)-7-[[[(6-Amino-2-pyridinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid;
 (S)-7-[[[(6-Ethylamino-2-pyridinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-
 10 methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid; and
 (±)-7-[[[(2-Amino-4-pyrimidinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid.

A preferred compound is (S)-7-[[[(6-Ethylamino-2-pyridinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid

- 15 Another embodiment of a benzodiazepine fibrinogen receptor template A is represented by the 1,4-benzodiazepine 2,5-dione of sub-formula (IV);



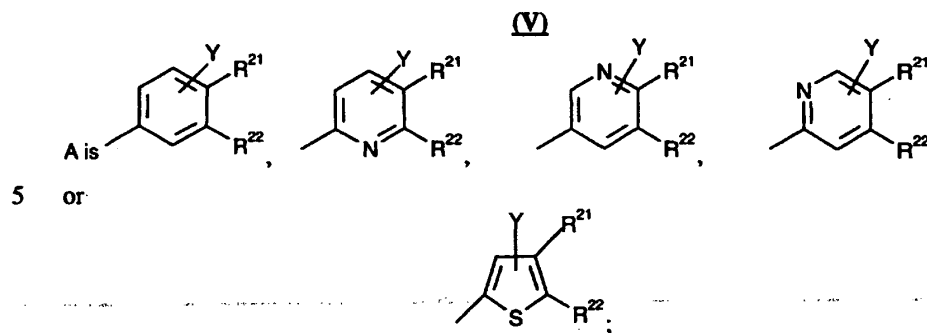
(IV)

wherein:

- Y is H, C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxycarbonyl, F, Cl, Br, I, CF₃, OR^f,
 20 S(O)_kR^f, COR^f, NO₂, N(R^f)₂, CO(NR^f)₂, CH₂N(R^f)₂, methylenedioxy, CN, CO₂R^f,
 OC(O)R^f, or NHC(O)R^f; and
 R^h is (CH₂)_qCO₂R^f.
 Suitably R^h is CH₂CH₂CO₂H.

- Entries (V)-(XV) in Table II summarize other illustrative fibrinogen receptor
 25 templates that are included within the scope of the present invention:

Table II



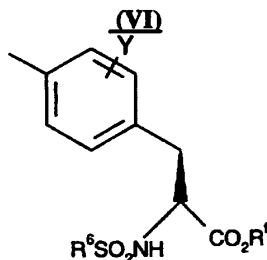
wherein:

R^{21} and R^{22} independently are H or $-Z-CO_2R^f$ or $Z-CON(R^f)_2$ with the proviso that one of R^{21} or R^{22} is $-Z-CO_2R^f$ or $Z-CON(R^f)_2$;

10 Z is $-CH_2-$, $-O(CH_2)_q-$, $-NR^f(CH_2)_q-$, $-S(CH_2)_q-$, $-CH_2CH_2-$, $-CH(CH_3)CH_2-$, $-(CH_2)_3-$, $-CH=CH-$, $-C(CH_3)=CH-$, $CH_2-CH=CH-$ or $CH=CHCH_2$; and

Y is H, C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-4} alkoxycarbonyl, F, Cl, Br, I, CF_3 , OR^f , $S(O)_kR^f$, COR^f , NO_2 , $N(R^f)_2$, $CO(NR^f)_2$, $CH_2N(R^f)_2$, methylenedioxy or $Z-COR^f$, disclosed in Alig, et al., EP 0 381 033, published August 8, 1990.

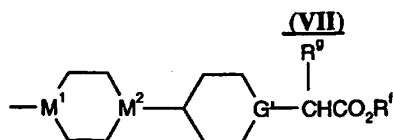
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wherein:

20 R^6 is aryl, C_{1-10} alkyl, C_{3-6} cycloalkyl, C_{4-10} aralkyl, C_{1-10} alkoxyalkyl, C_{1-10} alkaryl, C_{1-10} alkylthioalkyl, C_{1-10} alkoxythioalkyl, C_{1-10} alkylamino, C_{4-10} aralkylamino, C_{1-10} alkanoylamino, C_{4-10} aralkanoylamino, C_{1-10} alkanoyl, C_{4-10} aralkanoyl, or C_{1-10} carboxyalkyl; and

25 Y is H, C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-4} alkoxycarbonyl, F, Cl, Br, I, CF_3 , OR^f , $S(O)_kR^f$, COR^f , NO_2 , $N(R^f)_2$, $CO(NR^f)_2$, $CH_2N(R^f)_2$, methylenedioxy, CN, CO_2R^f , $OC(O)R^f$, or $NHC(O)R^f$, disclosed in Egbertson, et al., EP 0 478 328, published April 1, 1992.

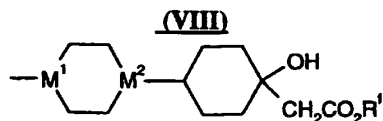


wherein:

- 5 M^1 is CH or N;
 M^2 is CH or N, with the proviso that when M^1 is CH, M^2 is N; and
 G' is N or $N^{\oplus}R''$,

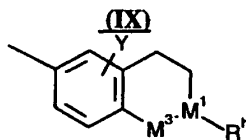
disclosed in Eldred, et al., EP 0542 363, published May 19, 1993.

10



wherein:

- M^1 is CH or N; and
 M^2 is CH or N, with the proviso that when M^1 is CH, M^2 is N,
 15 disclosed in Porter, et al., EP 0 537 980, published April 21, 1993.

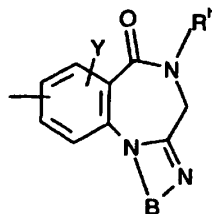


wherein:

- 20 M^1 is CH or N;
 Y is H, C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-4} alkoxycarbonyl, F, Cl, Br, I, CF_3 , OR^f ,
 $S(O)_kR^f$, COR^f , NO_2 , $N(R^f)_2$, $CO(NR^f)_2$, $CH_2N(R^f)_2$, methylenedioxy, CN, CO_2R^f ,
 $OC(O)R^f$, or $NHC(O)R^f$;
 D^3 is CH_2 or $C=O$; and
 25 R^h is $(CH_2)_qCO_2R^f$,

disclosed in Klinnick, et al., EP 0 635,492, published January 25, 1995.

(X)

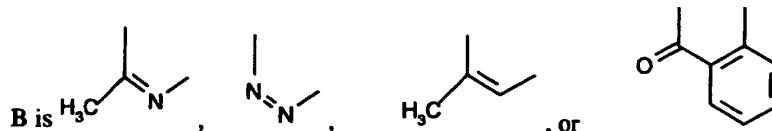


wherein:

Y is H, C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxycarbonyl, F, Cl, Br, I, CF₃, OR^f, S(O)_kR^f, COR^f, NO₂, N(R^f)₂, CO(NR^f)₂, CH₂N(R^f)₂, methylenedioxy, CN, CO₂R^f, OC(O)R^f, or NHC(O)R^f;

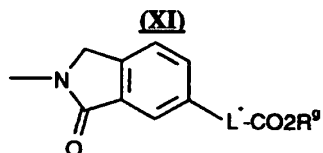
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R^h is (CH₂)_nCO₂R^f; and



disclosed in Blackburn, et al., WO 95/04057, published February 9, 1995.

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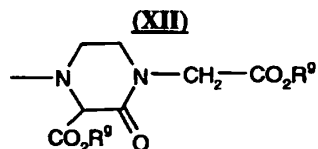


wherein:

L* is -C(O)NR^g-(CH₂)-, -C(O)-(CH₂)_q-, NR^g-(CH₂)_q-, -O-(CH₂)_q-, or S(O)_k-(CH₂)_q-;

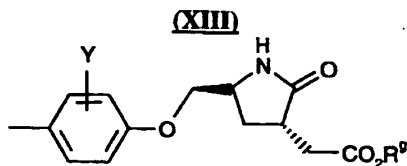
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disclosed in Hartman, et al., EP 0 540 331, published May 5, 1993.



disclosed in Sugihara, et al., EP 0 529,858, published March 3, 1993.

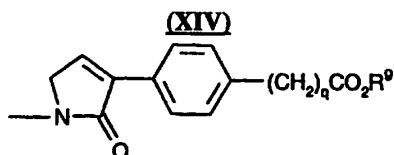
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wherein:

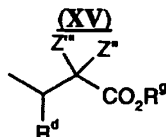
Y is H, C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkoxycarbonyl, F, Cl, Br, I, CF₃, OR^f, S(O)_kR^f, COR^f, NO₂, N(R^f)₂, CO(NR^f)₂, CH₂N(R^f)₂, methylenedioxy, CN, CO₂R^f, OC(O)R^f, or NHC(O)R^f, disclosed in Himmeisbach, et al., EP 0 483 667, published May 6, 1992.

5



disclosed in Linz, et al., EP 0 567 968, published November 3, 1993.

10



wherein:

R^d is Het-C₀₋₆alkyl; and

Z'', Z''' independently are hydrogen, C₁₋₄alkyl, halo, OR^f, CN, S(O)_kR^f,

15 CO₂R^f, or OH,

disclosed in Bovy, et al., EP 0 539 343, published April 28, 1993.

The above descriptions of fibrinogen receptor templates for use in the present invention were taken from pending published patent applications. Reference should be made to such patent applications for their full disclosures, including the variations possible for such templates and methods of preparing said templates, the entire disclosure of such patent applications being incorporated herein by reference.

In cases wherein the compounds of this invention may have one or more chiral centers, unless specified, this invention includes each unique nonracemic compound which may be synthesized and resolved by conventional techniques. In cases in which compounds have unsaturated carbon-carbon double bonds, both the cis (Z) and trans (E) isomers are within the scope of this invention. The meaning of any substituent at any one occurrence is independent of its meaning, or any other substituent's meaning, at any other occurrence.

Abbreviations and symbols commonly used in the peptide and chemical arts are used herein to describe the compounds of this invention. In general, the amino acid abbreviations follow the IUPAC-IUB Joint Commission on Biochemical Nomenclature as described in *Eur. J. Biochem.*, 158, 9 (1984).

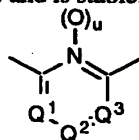
C₁₋₄alkyl as applied herein means an optionally substituted alkyl group of 1 to 4 carbon atoms, and includes methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl and t-butyl. C₁₋₆alkyl additionally includes pentyl, n-pentyl, isopentyl, neopentyl and hexyl and the simple aliphatic isomers thereof. C₀₋₄alkyl and C₀₋₆alkyl additionally indicates
 5 that no alkyl group need be present (e.g., that a covalent bond is present).

Any C₁₋₄alkyl or C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl or C₁₋₆ oxoalkyl may be optionally substituted with the group R^x, which may be on any carbon atom that results in a stable structure and is available by conventional synthetic techniques. Suitable groups for R^x are C₁₋₄alkyl, OR', SR', C₁₋₄alkyl, C₁₋₄alkylsulfonyl, C₁₋₄alkylsulfoxyl, -CN,
 10 N(R')₂, CH₂N(R')₂, -NO₂, -CF₃, -CO₂R', -CON(R')₂, -CO R', -N R'C(O) R', OH, F, Cl, Br, I, N₃ or CF₃S(O)_r, wherein r is 0 to 2 and R' is as defined with respect to formula (II).

Ar, or aryl, as applied herein, means phenyl or naphthyl, or phenyl or naphthyl substituted by one to three substituents, such as those defined above for alkyl, especially C₁₋₄alkyl, C₁₋₄alkoxy, C₁₋₄alkthio, CO₂H, N₃, trifluoroalkyl, OH, F, Cl, Br or I.

15 Het, or heterocycle, indicates an optionally substituted five or six membered monocyclic ring, or a nine or ten-membered bicyclic ring containing one to three heteroatoms chosen from the group of nitrogen, oxygen and sulfur, which are stable and available by conventional chemical synthesis. Illustrative heterocycles are benzofuryl, benzimidazole, benzopyran, benzothiophene, biotin, furan, imidazole, indoline,
 20 morpholine, piperidine, piperazine, pyrrole, pyrrolidine, tetrahydropyridine, pyridine, thiazole, thiophene, quinoline, isoquinoline, and tetra- and perhydro- quinoline and isoquinoline. Any accessible combination of up to three substituents on the Het ring, such as those defined above for alkyl that are available by chemical synthesis and are stable are within the scope of this invention.

25 C₃₋₇cycloalkyl refers to an optionally substituted carbocyclic system of three to seven carbon atoms, which may contain up to two unsaturated carbon-carbon bonds. Typical of C₃₋₇cycloalkyl are cyclopropyl, cyclobutyl, cyclopentyl, cyclopentenyl, cyclohexyl, cyclohexenyl and cycloheptyl. Any combination of up to three substituents, such as those defined above for alkyl, on the cycloalkyl ring that is available by
 30 conventional chemical synthesis and is stable, is within the scope of this invention.



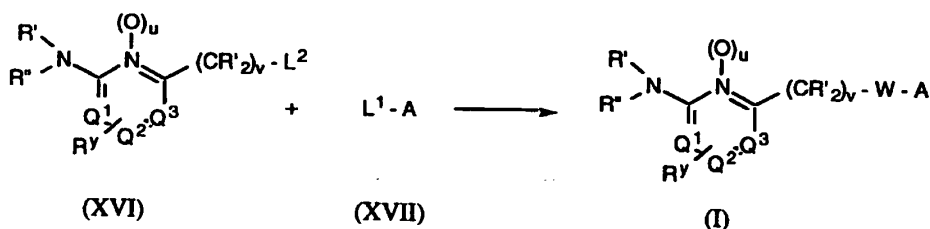
The ring represented by is a six-membered ring containing at least one nitrogen which is 2,6-disubstituted relative to such nitrogen. The ring may optionally have an additional nitrogen atom in the ring, and hence may be a pyrazine or a pyrimidine. The substituent R^y may be in any position on Q¹ - Q³ which results in a

stable structure. It will be apparent that when the value of u is 1 the compound described will be an N-oxide; whereas, when the value of u is 0 there is no oxygen substituent on the nitrogen. A pyridine ring is preferred.

Certain radical groups are abbreviated herein. t -Bu refers to the tertiary butyl radical, Boc refers to the t -butoxycarbonyl radical, Fmoc refers to the fluorenylmethoxycarbonyl radical, Ph refers to the phenyl radical, Cbz refers to the benzyloxycarbonyl radical, BrZ refers to the o -bromobenzyloxycarbonyl radical, ClZ refers to the o -chlorobenzyloxycarbonyl radical, Bzl refers to the benzyl radical, 4-MBzl refers to the 4-methyl benzyl radical, Me refers to methyl, Et refers to ethyl, Ac refers to acetyl, Alk refers to C_{1-4} alkyl, Nph refers to 1- or 2-naphthyl and cHex refers to cyclohexyl. Tet refers to 5-tetrazolyl.

Certain reagents are abbreviated herein. DCC refers to dicyclohexylcarbodiimide, DMAP refers to dimethylaminopyridine, DIEA refers to diisopropylethyl amine, EDC refers to 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide, hydrochloride. HOBt refers to 1-hydroxybenzotriazole, THF refers to tetrahydrofuran, DIEA refers to diisopropylethylamine, DME refers to dimethoxyethane, DMF refers to dimethylformamide, NBS refers to N -bromosuccinimide, Pd/C refers to a palladium on carbon catalyst, PPA refers to 1-propanephosphonic acid cyclic anhydride, DPPA refers to diphenylphosphoryl azide, BOP refers to benzotriazol-1-yloxy-tris(dimethyl-amino)phosphonium hexafluorophosphate, HF refers to hydrofluoric acid, TEA refers to triethylamine, TFA refers to trifluoroacetic acid, PCC refers to pyridinium chlorochromate.

Compounds of the formula (I) are generally prepared by reacting a compound of formula (XVI) with a compound of formula (XVII), wherein L^1 and L^2 are groups which may react to form a covalent bond in the moiety W, by methods generally known in the art.



Typical methods include coupling to form amide bonds, nucleophilic displacement reactions and palladium catalyzed couplings.

For instance, when W contains an ether or amine linkage, the bond may be formed by a displacement reaction, and one of L^1 and L^2 will contain an amino or hydroxy group

and the other will contain a displaceable group, such as a chloro, bromo or iodo group.

When W contains an amide bond, typically one of L¹ and L² will contain an amino group, and the other will contain a carboxylic acid group. In another approach, L¹ may be an aryl or heteroaryl bromide, iodide or trifluoromethylsulfonyloxy derivative and L² may contain an amino group and the amide linkage may be formed by palladium-catalyzed aminocarbonylation with carbon monoxide in a suitable solvent such as dimethylformamide or toluene.

It will be apparent that the precise identity of L¹ and L² will be dependent upon the site of the linkage being formed. General methods for preparing the linkage

(CHR")_r-U-(CHR")_s-V- are described, for example, in EP-A 0 372 486 and EP-A 0 381 033 and EP-A 0 478 363, which are incorporated herein by reference.

For instance, if V is CONH, L¹ may be -NH₂, L² may be OH (as in an acid) or Cl (as in an acid chloride). For instance, (2-amino-pyrid-6-yl)(CH₂)_a-COCl may be reacted with a suitable amine. When L² is OH, a coupling agent is used.

Similarly, if V is NHCO, L¹ may be -CO₂H or CO-Cl, L² may be -NH₂. For instance, (2-amino-pyrid-6-yl)(CH₂)_a-NHR' may be reacted with a suitable carboxylic acid.

Where V is NHSO₂, L¹ may be SO₂Cl, L² may be -NH₂ as above. Where V is SO₂NH, L¹ may be -NH₂ and L² may be SO₂Cl. Methods to prepare such sulfonyl chlorides are disclosed, for instance, in *J. Org. Chem.*, 23, 1257 (1958).

If V is CH=CH, L¹ may be -CHO, L² may be CH=P-Ph₃. Alternately, L¹ may be CH=P-Ph₃ and L² may be CHO. For instance, (2-amino-pyrid-6-yl)(CH₂)_a-CHO may be reacted with a suitable phosphorane.

Where V is CH₂CH₂ may be obtained by reduction of a suitably protected compound wherein V is CH=CH.

Where V is CH₂O, CH₂N or C≡C, L¹ may be -OH, -NH or -C≡CH, respectively; L² may be -Br or -I. Similarly where U or V is OCH₂, NR'CH₂ or C≡C, L¹ may be -CH₂Br and L² may be -OH, -NH or -C≡CH, respectively. For example, (2-amino-pyrid-6-yl)(CH₂)_a-Br may be reacted with an appropriate amine, alkoxide or acetylene.

Alternately, when U or V is $C\equiv C$, L^1 may be Br, I or CF_3SO_3 , L^2 may be $C\equiv CH$ and the coupling may be catalyzed by palladium and a base.

Compounds wherein V is $CHOHCH_2$ may be prepared from a suitably protected compound where V is $CH=CH$ by the procedure disclosed in *J. Org. Chem.*, 54, 1354

5 (1989).

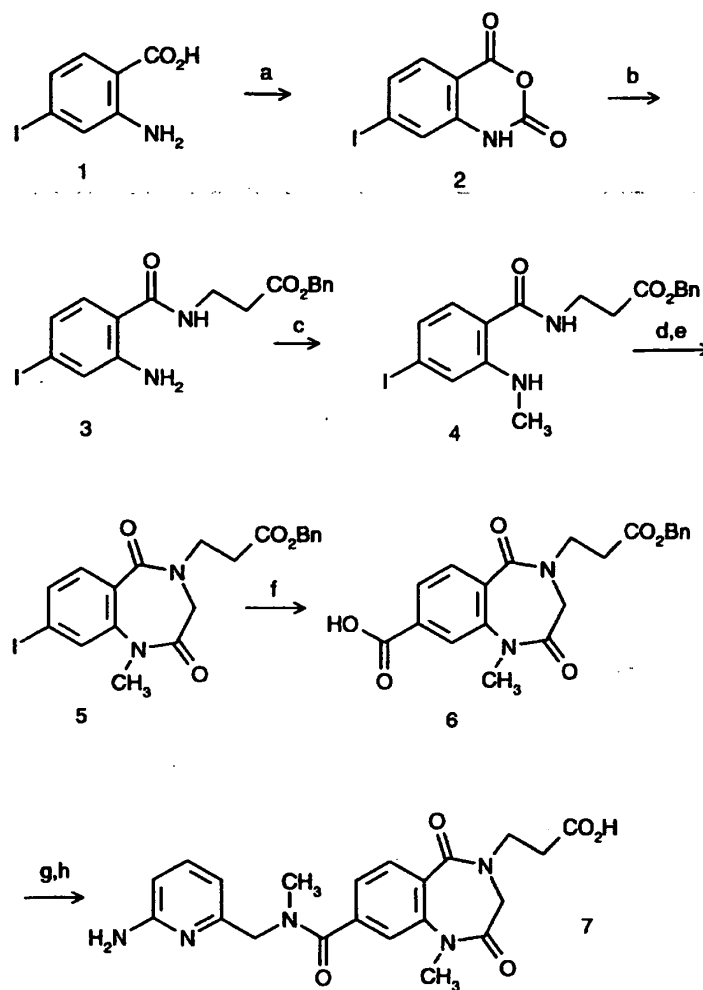
Compounds wherein V is CH_2CHOH may be obtained from a suitably protected compound where V is $CH=CH$ by hydroboration and basic oxidation as disclosed in *Tet. Lett.*, 31, 231 (1990).

The core 6-7 fused ring fibrinogen template of formula (II) is prepared by methods
10 well known in the art, e.g., Hynes, *et al.*, *J. Het. Chem.*, 1988, 25, 1173; Muller, *et al.*, *Helv. Chim. Acta.*, 1982, 65, 2118; Mori, *et al.*, *Heterocycles*, 1981, 16, 1491. Similarly, methods for preparing benzazepines, 1,4-benzothiazepines, 1,4-benzoxazepines and 1,4-benzodiazepines are known and are disclosed, for instance, in Bondinell, *et al.*, International Patent Application WO 93/00095.

15 Representative fibrinogen antagonist templates may be prepared according to Schemes A - AA, which follow:

Scheme A describes a method of exemplary fibrinogen receptor templates described in Blackburn, et. al., WO 93/08174.

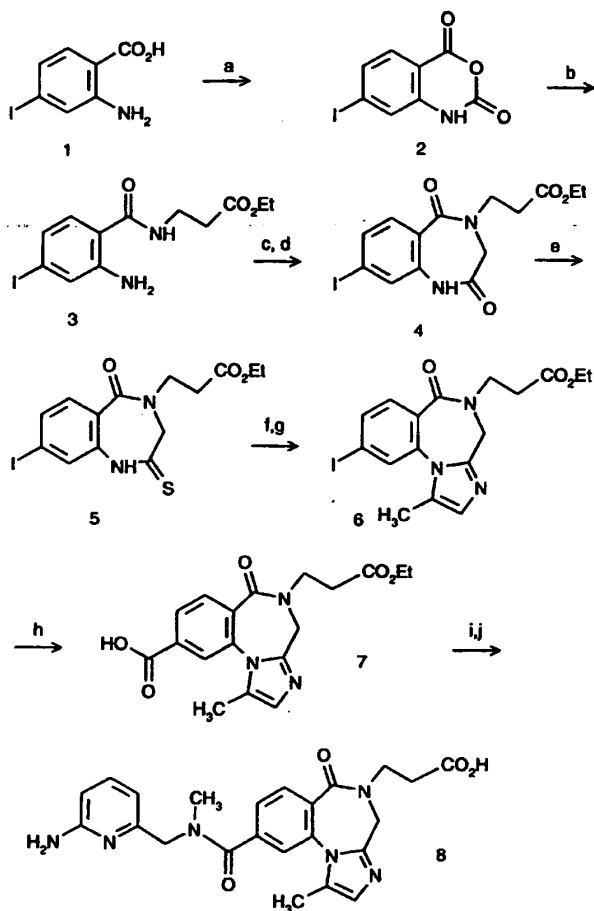
Scheme A



- a) COCl_2 , Na_2CO_3 , toluene; b) β -alanine benzyl ester tosylate, DMAP, pyridine; c) CH_3I , 2,6-lutidine, DMF; d) α -bromoacetyl bromide, Et_3N , CH_2Cl_2 ; e) NaH , DMF, f) $\text{Pd}(\text{OAc})_2$, dppf, CO , DMSO, 65°C , 18 h; g) 6-(methylamino)methyl-2-pyridinamine, EDC, HOBT· H_2O , DIEA, CH_3CN ; h) H_2 , 10% Pd/C , EtOH .

Scheme B describes a method of preparing exemplary fibrinogen receptor templates described in Blackburn, *et. al.*, WO 95/04057.

Scheme B

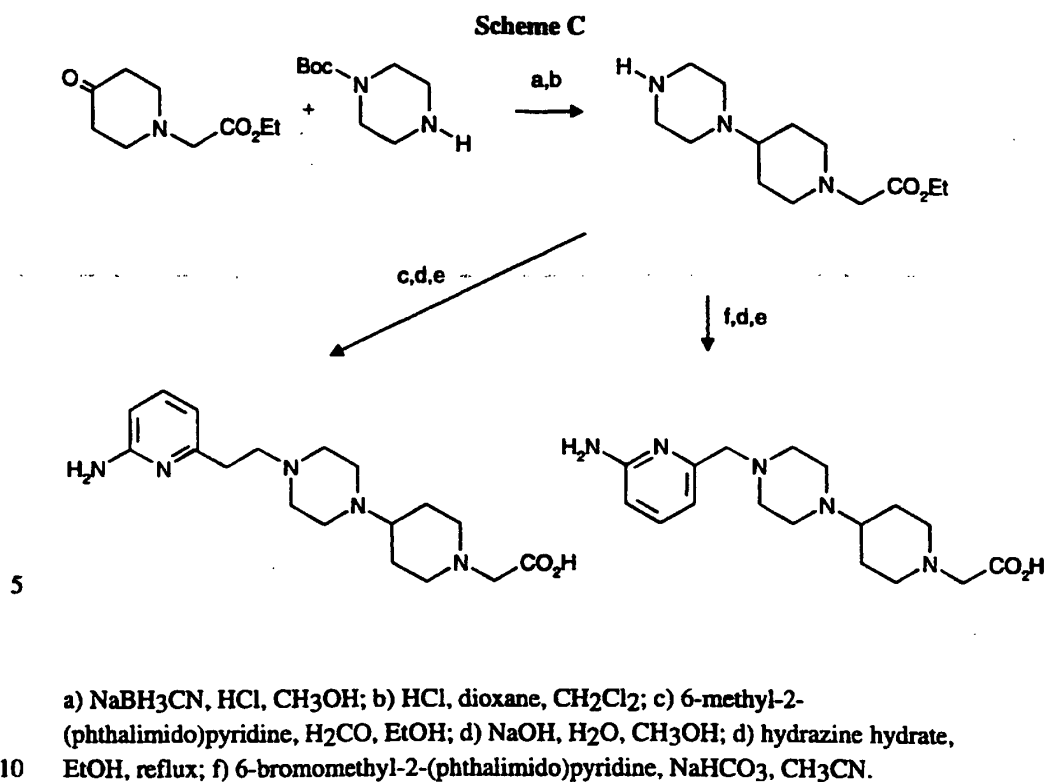


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- a) COCl_2 , NaHCO_3 , toluene; b) β -alanine ethyl ester hydrochloride, DMAP, pyridine; c) α -bromoacetyl bromide, Et_3N , CH_2Cl_2 ; d) NaH , DMF; e) Lawesson's reagent, THF, 50°C , 2 h; f) CH_3I , NaOH , $(n\text{-Bu})_4\text{N}\cdot\text{HSO}_4$, CH_2Cl_2 , H_2O , RT, 2 h; g) propargyl amine, toluene, pyridine hydrochloride, reflux, 6 h; h) $\text{Pd}(\text{OAc})_2$, dppf, CO, DMSO, 65°C , 18 h; i) 6-(methylamino)methyl-2-pyridinamine, EDC, HOBT· H_2O , DIEA, CH_3CN ; j) LiOH , H_2O , THF, 18 h.

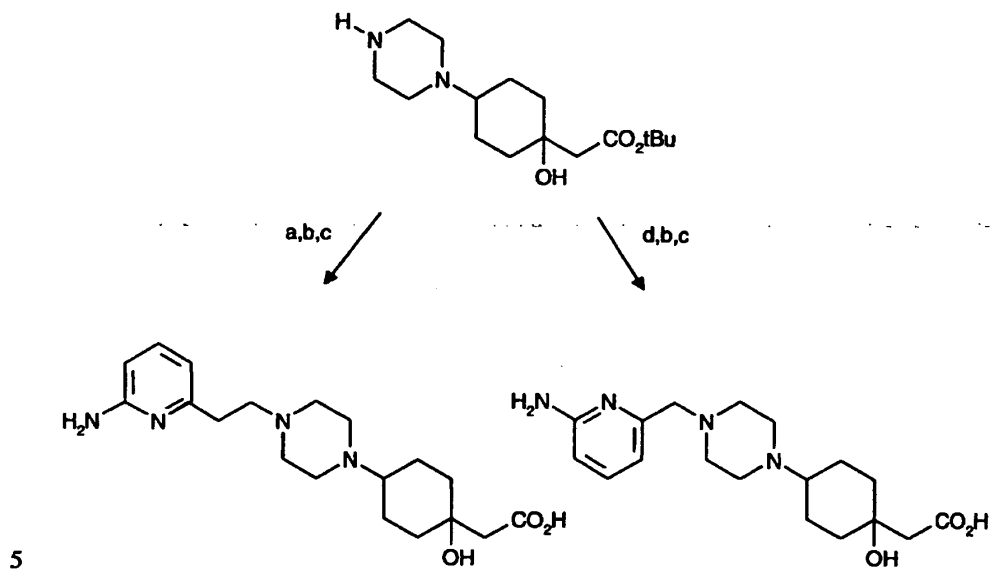
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Scheme C describes a method of preparing exemplary fibrinogen receptor templates described in Porter, *et al.*, EP 0542363.



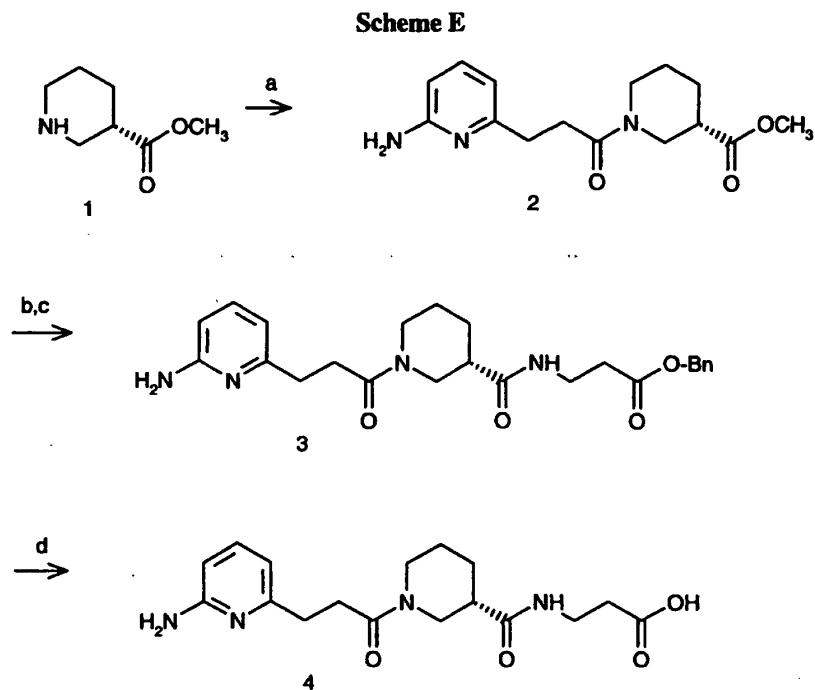
Scheme D describes a method of preparing exemplary fibrinogen receptor templates described in Porter, *et al.*, EP 0537980.

Scheme D



a) 6-methyl-2-(phthalimido)pyridine, H_2CO , EtOH; b) NaOH, H_2O , CH_3OH ; c) hydrazine hydrate, EtOH, reflux; d) 6-bromomethyl-2-(phthalimido)pyridine, NaHCO_3 , CH_3CN .

Scheme E describes a method of preparing exemplary fibrinogen receptor templates described in Beavers, *et al.*, WO 95/25091.



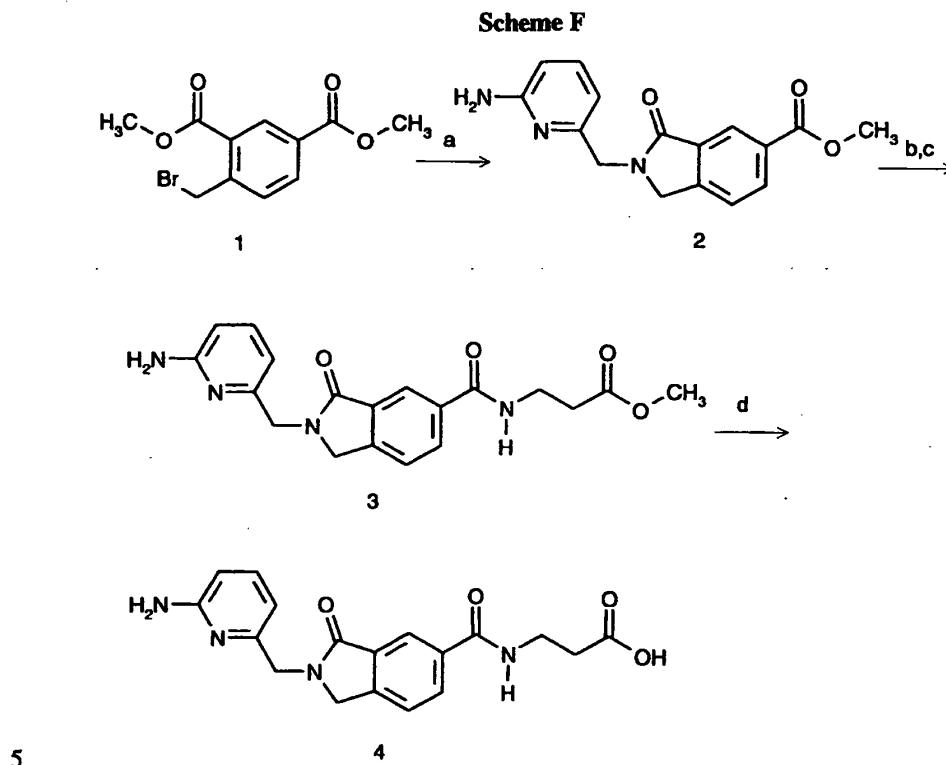
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a) (6-amino-2-pyridinyl)propionic acid, BOP-Cl, NMM, CH₂Cl₂; b) LiOH, H₂O, THF;
c) β-alanine benzyl ester, EDC, HOBT, NMM, CH₂Cl₂; d) H₂, 10% Pd/C, AcOH, THF,
H₂O.

10

Following the procedures of Beavers, *et al.*, WO 95/25091, Example 1, except substituting 3-(6-amino-2-pyridinyl)propionic acid, Bondinell, *et al.*, WO 95/25091, for N^α-Boc-D-lys(Cbz)-OH, gives E-4.

Scheme F describes a method of preparing exemplary fibrinogen receptor templates described in Hartman, *et al.*, EP 0540334.

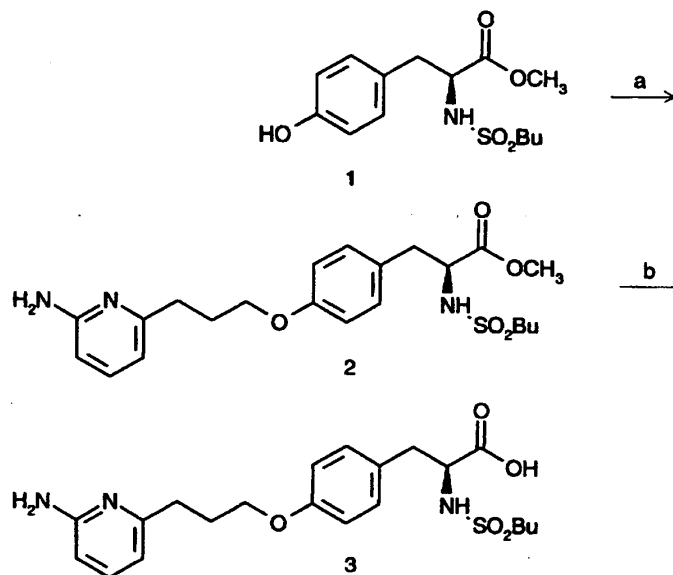


a) 6-aminomethyl-2-pyridinamine, Et₃N, benzene; b) 1.0 N LiOH, H₂O, CH₃OH; c) β-alanine ethyl ester, BOP, Et₃N, CH₃CN; d) LiOH, H₂O, THF, CH₃OH.

- 10 Dimethyl 4-(bromomethyl)benzene-1,3-dicarboxylate, **F-1**, is treated with a suitably functionalized amine, such as 6-aminomethyl-2-pyridinamine, under the general conditions described for 2,3-dihydro-N-(2-carboxy-ethyl)-2-[2-(piperidinyl)ethyl]-3-oxo-1H-isoindole-5-carboxamide in Hartman, *et al.*, EP 0540334, to give **F-4**.

Scheme G describes a method of preparing exemplary fibrinogen receptor templates described in Egbertson, *et al.*, EP 0478363.

Scheme G



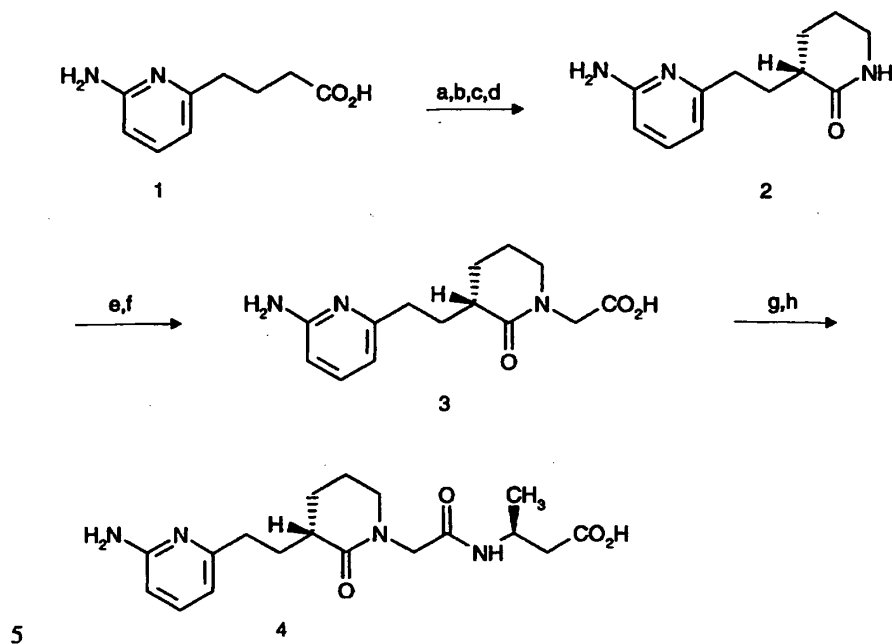
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a) 3-(6-amino-2-pyridinyl)propanol, Ph_3P , DEAD, CH_2Cl_2 , benzene; b) 1.0 N LiOH, THF, H_2O .

- 10 N-(n-Butylsulfonyl)-L-tyrosine methyl ester, **G-1**, is treated with a suitably functionalized alcohol, such as 3-(6-amino-2-pyridinyl)propanol, prepared following the procedures of Warter, *et al.*, *Org. Synth.* **1943**, 23, 83, and Bruckelman, *et al.*, *J. Chem. Soc. Perkin Trans. I*, **1984**, 2801-2807, to give **G-3**.

Scheme H describes a method of preparing exemplary fibrinogen receptor templates described in Duggan, *et al.*, *J. Med. Chem.* **1995**, *38*, 3332.

Scheme H



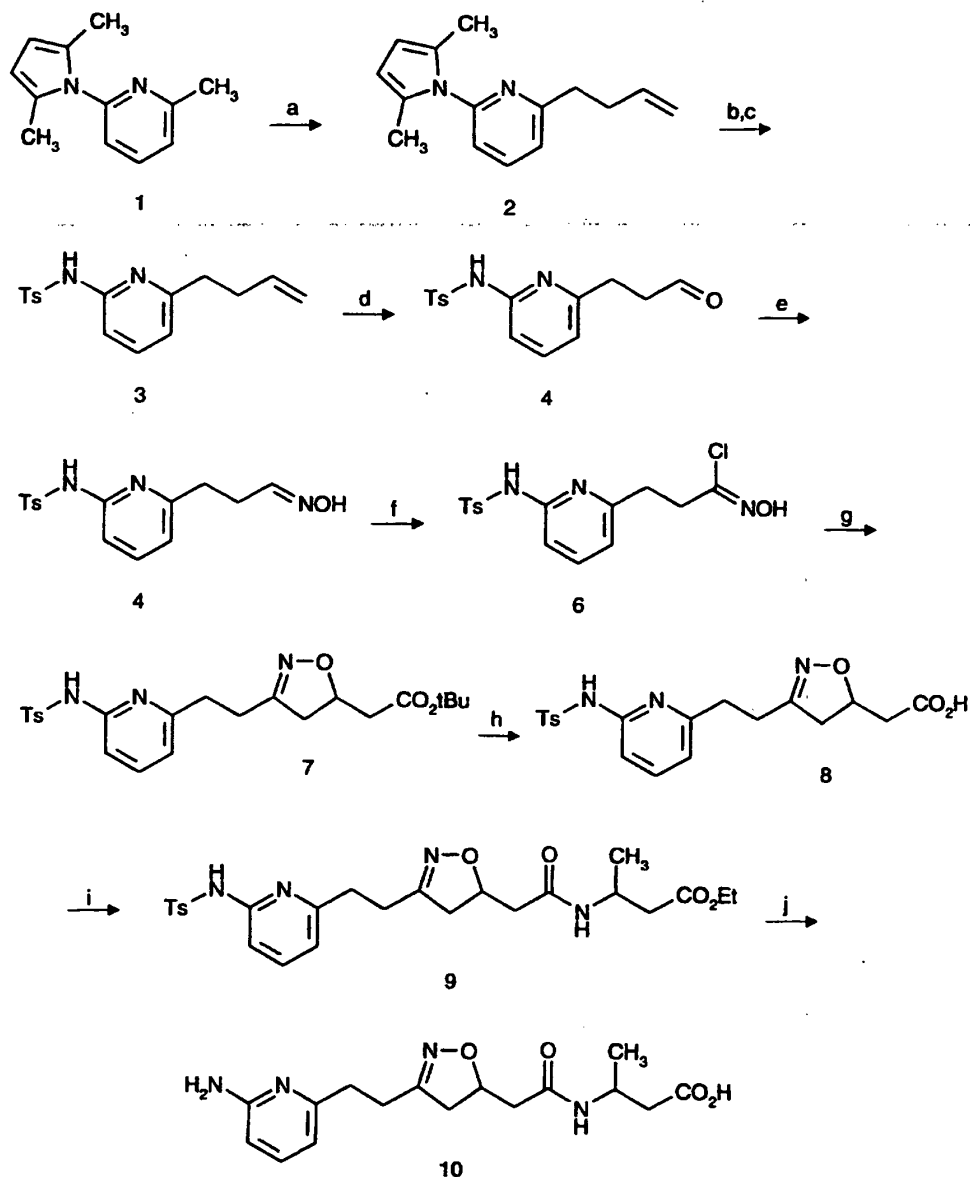
- a) pivaloyl chloride, Et₃N, THF, (S)-4-benzyl-2-oxazolidinone; b) Ti(O-*i*-Pr)₂, acrylonitrile, DIEA, CH₂Cl₂; c) H₂, PtO₂, CH₃OH, CHCl₃; d) NaHCO₃, CH₃CN; e) NaHMDS, ethyl bromoacetate; f) 1 N NaOH, CH₃OH; g) 3(R)-methyl-β-alanine ethyl ester HCl, EDC, HOBT, Et₃N, DMF; h) 1 N NaOH, CH₃OH.

A suitably functionalized carboxylic acid, such as 4-(6-amino-2-pyridinyl)butanoic acid, **H-1**, is activated and reacted with a chiral auxiliary such as lithium (S)- 4-benzyl-2-oxazolidinone to form a chiral Evans reagent. Alkylation of the titanium enolate with acrylonitrile, followed by nitrile reduction and lactam formation affords lactam **H-2**. Alkylation of the lactam with agents such as ethyl bromoacetate followed by ester saponification yields the carboxylic acid **H-3**. The resulting carboxylic acid derivative **H-3** is converted to an activated form of the carboxylic acid using, for example, EDC and HOBT, or SOCl₂, and the activated form is subsequently reacted with

- an appropriate amine, for instance the 3(R)-methyl- β -alanine ethyl ester, in a suitable solvent such as DMF, CH_2Cl_2 , or CH_3CN . Depending on whether acid neutralization is required, an added base, such as DIEA or pyridine, may be used. Many additional methods for converting a carboxylic acid to an amide are known, and can be found in
- 5 standard reference books, such as "Compendium of Organic Synthetic Methods", Vol. I - VI (published by Wiley-Interscience), or Bodansky, "The Practice of Peptide Synthesis" (published by Springer-Verlag). Hydrolysis of the ethyl ester is accomplished according to the general conditions described for the conversion of H-2 to H-3, to provide the carboxylic acid H-4. Alternatively, the intermediate carboxylate salt of can be isolated, if
- 10 desired, or a carboxylate salt of the free carboxylic acid can be prepared by methods well-known to those of skill in the art.

Scheme I describes a method of preparing exemplary fibrinogen receptor templates described in WO 93/07867.

Scheme I



a) LDA, THF, allyl bromide; b) $\text{NH}_2\text{OH}\cdot\text{HCl}$, EtOH, H_2O ; c) TsCl , NaH, THF; d) O_3 , CH_2Cl_2 , CH_3OH , DMS; e) $\text{NH}_2\text{OH}\cdot\text{HCl}$, NaOAc, CH_3OH ; f) NCS, DMF; g) tert-butyl 3-butenate, Et_3N ; h) 4M HCl, dioxane, CH_2Cl_2 ; i) ethyl 3-aminobutyrate, EDC, $\text{HOBT}\cdot\text{H}_2\text{O}$, DIEA, CH_3CN ; j) 1.0 N LiOH, THF, H_2O .

5

The readily available aminopyridine derivative **I-1**, *J. Chem. Soc. Perkin Trans. I* **1984**, 2801, is converted to the alkylated derivative **I-2** by the general protocol described by Meakins, *J. Chem. Soc. Perkin Trans. I* **1984**, 2801. Thus, **I-1** is deprotonated with an amide base, such as lithium diisopropylamide or lithium bis(trimethylsilyl)amide, and the
10 resulting metalated species is alkylated with an appropriate alkylating agent, for instance allyl bromide, to afford the butenyl derivative **I-2**. Generally, THF or ethylene glycol dimethyl ether are the solvents of choice for an alkylation reaction, although THF in the presence of various additives, for instance HMPA or TMEDA, can be used. The 2,5-dimethylpyrrole protecting group is conveniently removed at this stage using the general
15 protocol described by Meakins (see reference above). Thus, **I-2** is reacted with hydroxylamine hydrochloride in an appropriate solvent, e.g., aqueous EtOH, to afford the corresponding deprotected aminopyridine. Protection of the amino group of this aminopyridine can be accomplished by reaction with a sulfonyl chloride, for instance p-toluenesulfonyl chloride, in the presence of a suitable base, generally NaH or an aqueous
20 alkali metal hydroxide, in an inert solvent, preferably THF, to afford **I-3**. Alternative protecting groups known to those of skill in the art may be used, as long as they are compatible with the subsequent chemistry and can be removed when desired. Such protecting groups are described in Greene, "Protective Groups in Organic Synthesis" (published by Wiley-Interscience). Oxidative cleavage of the olefin of **I-3** to afford the
25 aldehyde **I-4** can be conveniently accomplished by ozonolysis in an inert solvent, usually CH_2Cl_2 or a mixture of CH_2Cl_2 and CH_3OH , followed by in-situ reduction of the ozonide with a suitable reducing agent, generally dimethylsulfide (DMS) or triphenylphosphine. Alternative methods for oxidative cleavage, such as the Lemieux-Johnson reaction, *J. Org. Chem.* **1956**, 21, 478, can also be used. The aldehyde is
30 converted to the aldoxime **I-5** by standard procedures known to those of skill in the art, and this aldoxime is oxidized to the oximinoyl chloride derivative **I-6** by the methods described in WO 95/14682 and WO 95/14683. Reaction of **I-6** with an olefin, such as

tert-butyl 3-butenate, *Tet. Lett.* **1985**, 26, 381-384, in the presence of a suitable base, for instance Et₃N or DIEA, in an inert solvent such as benzene or toluene, according to the protocol described in WO 95/14682 and WO 95/14683, gives the cycloadduct **I-7**. The tert-butyl ester of **I-7** is removed under standard acidic conditions, generally TFA in

5 CH₂Cl₂ or HCl in dioxane, to give the carboxylic acid **I-8**. The carboxylic acid is activated using, for example, EDC and HOBt, or SOCl₂, and the activated form is subsequently reacted with an appropriate amine, for instance a suitable derivative of β-alanine, in a neutral solvent, such as DMF, CH₂Cl₂, or CH₃CN, to afford **I-9**. Depending

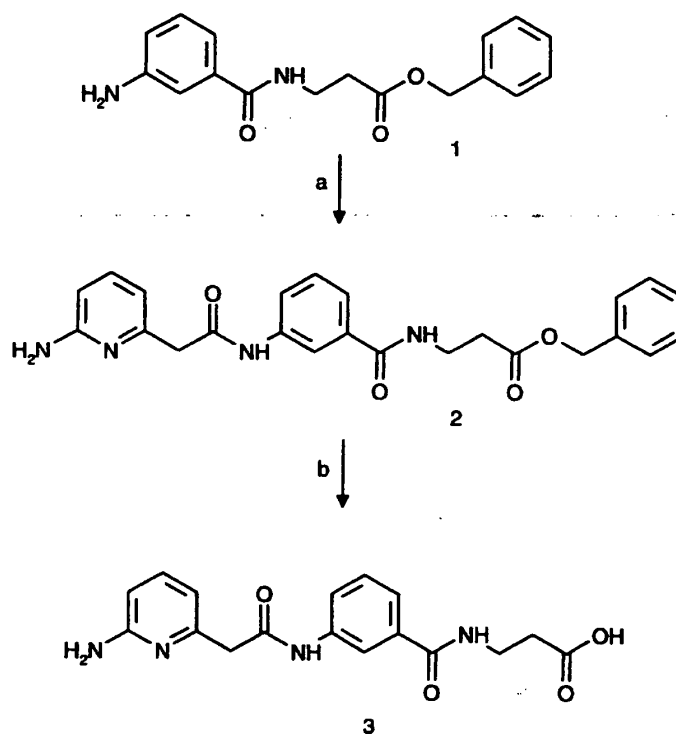
10 on whether acid neutralization is required, an added base, such as DIEA or pyridine, may be used. Many additional methods for converting a carboxylic acid to an amide are known, and can be found in standard reference books, such as "Compendium of Organic Synthetic Methods", Vol. I - VI (published by Wiley-Interscience), or Bodansky, "The Practice of Peptide Synthesis" (published by Springer-Verlag). Derivatives of β-alanine

15 are readily available in either racemic or optically pure form by a variety of methods known to those of skill in the art. A representative method is described in WO 93/07867. The ethyl ester and sulfonyl protecting groups of **I-9** are removed using aqueous base, for example, LiOH in aqueous THF or NaOH in aqueous CH₃OH or EtOH. The intermediate carboxylate salt is acidified with a suitable acid, for instance TFA or HCl, to afford the carboxylic acid **I-10**. Alternatively, the intermediate carboxylate salt can be

20 isolated, if desired, or a carboxylate salt of the free carboxylic acid can be prepared by methods well-known to those of skill in the art.

Scheme J describes a method of preparing exemplary fibrinogen receptor templates described in Alig, *et al.*, EP 0372486.

Scheme J

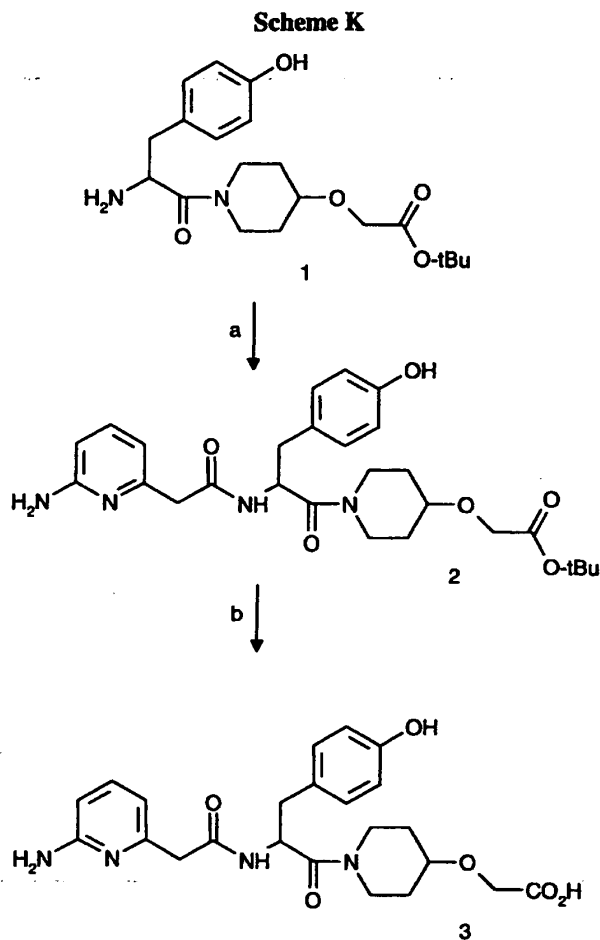


a) (6-amino-2-pyridinyl)acetic acid, EDC, DIEA, DMF; b) NaOH, H₂O, CH₃OH.

J-1, prepared as described in Alig *et al.*, EP 0372486, is condensed with a suitable substituted carboxylic acid, such as (6-amino-2-pyridinyl)acetic acid, prepared by saponification of ethyl (6-amino-2-pyridinyl)acetate, Awaya, *et al.*, *Chem. Pharm. Bull.* 1974, 22, 1414, in the presence of EDC and DIEA, and in a suitable solvent, e.g., DMF or CH₃CN, to give J-2. Many additional methods for converting a carboxylic acid to an amide are known, and can be found in standard reference books, such a "Compendium of Organic Synthesis", Vol. I-VI (published by Springer-Verlag). Hydrolysis of the ester in

J-2 is accomplished by saponification with a suitable reagent, e.g., NaOH, in a suitable solvent, e.g., aqueous methanol. Alternatively, the benzyl ester in J-2 may be converted to the acid by treatment with hydrogen and a suitable catalyst, e.g., Pd/C, in a suitable solvent, e.g., CH₃OH, EtOH, or AcOH.

- 5 Scheme K describes a method of preparing exemplary fibrinogen receptor templates described in Alig, *et al.*, EP 0505868.



10

a) (6-amino-2-pyridinyl)acetic acid, EDC, DIEA, DMF; b) CF₃CO₂H, CH₂Cl₂.

K-1, prepared as described in Alig et al., EP 0505868, is condensed with a suitable substituted carboxylic acid, such (6-amino-2-pyridinyl)acetic acid, prepared by saponification of ethyl (6-amino-2-pyridinyl)acetate, Awaya, *et al.*, *Chem. Pharm. Bull.* **1974**, *22*, 1414, in the presence of EDC and DIEA, in a suitable solvent, e.g., DMF or

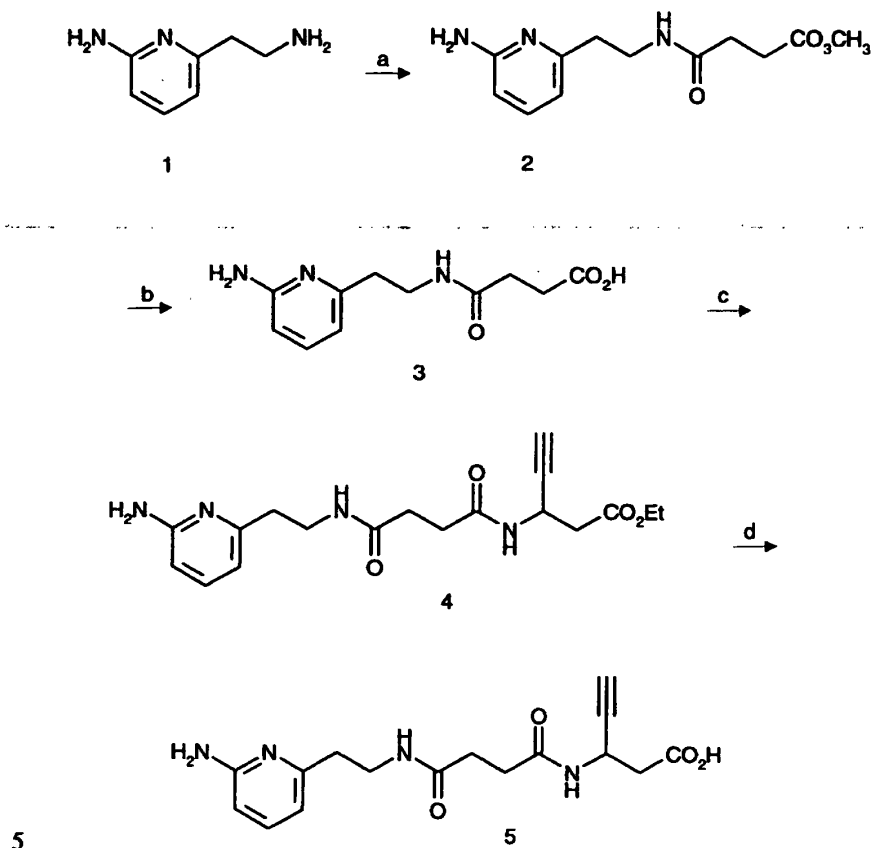
5 **CH₃CN**, to give **K-2**. Many additional methods for converting a carboxylic acid to an amide are known, and can be found in standard reference books, such as "Compendium of Organic Synthesis", Vol. I-VI (published by Springer-Verlag). Hydrolysis of the ester in **J-2** is accomplished with trifluoroacetic acid or hydrogen chloride to give **K-3**.

Alternatively, the ester in **K-2** may be saponified with a suitable reagent, e.g., 1N NaOH,

10 in a suitable solvent, e.g., **CH₃OH**.

Scheme L describes a method of preparing exemplary fibrinogen receptor templates described in WO 93/07867.

Scheme L



a) 3-(carbomethoxy)propionyl chloride, DIEA, CH_2Cl_2 ; b) 1.0 N NaOH, CH_3OH ; c) ethyl 3-amino-4-pentynoate, EDC, HOBt· H_2O , DIEA, CH_3CN ; d) 1.0 N LiOH, THF, H_2O .

10

A suitably functionalized amine, such as 2-amino-6-(2-aminoethyl)pyridine, prepared following the procedures of Preparation 13 in Bondinell, *et al.*, WO 93/00095, for the preparation of 2-aminopyridine-4-ethanamine dihydrochloride, except substituting 2-amino-6-picoline for the 2-amino-4-picoline is reacted with 3-(carbomethoxy)propionyl

chloride in the presence of an appropriate acid scavenger, such as Et₃N, DIEA, or pyridine, in a neutral solvent, generally CH₂Cl₂, to afford L-2. The methyl ester of L-2 is hydrolyzed using aqueous base, for example, LiOH in aqueous THF or NaOH in aqueous CH₃OH or EtOH, and the intermediate carboxylate salt is acidified with a

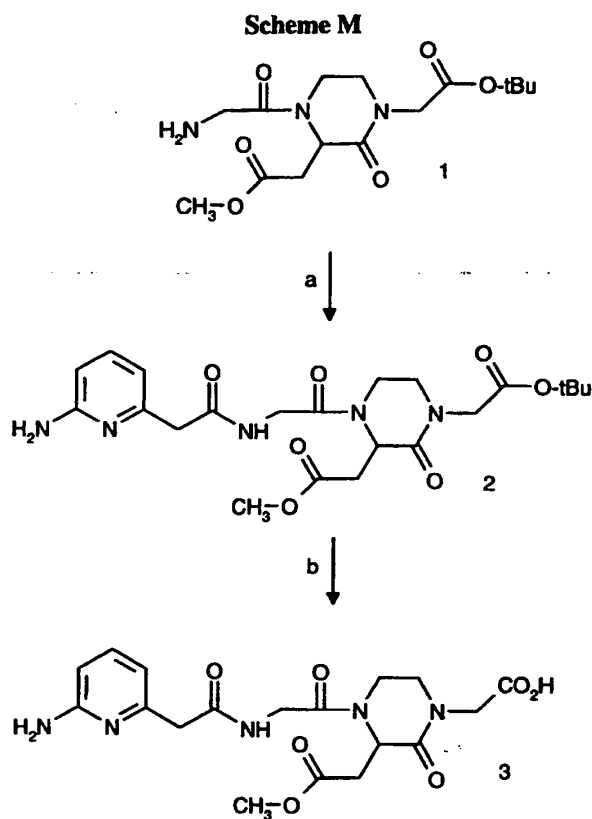
5 suitable acid, for instance TFA or HCl, to afford the carboxylic acid L-3. Alternatively, L-1 can be reacted with succinic anhydride in the presence of an appropriate base, such as Et₃N, DIEA, or pyridine, in a neutral solvent, generally CH₂Cl₂, to afford L-3 directly. The resulting carboxylic acid derivative L-3 is converted to an activated form of the carboxylic acid using, for example, EDC and HOBt, or SOCl₂, and the activated form is

10 subsequently reacted with an appropriate amine, for instance the known ethyl 3-amino-4-pentynoate (WO 93/07867), in a suitable solvent such as DMF, CH₂Cl₂, or CH₃CN, to L-4. Depending on whether acid neutralization is required, an added base, such as DIEA or pyridine, may be used. Many additional methods for converting a carboxylic acid to an amide are known, and can be found in standard reference books, such as "Compendium of

15 Organic Synthetic Methods", Vol. I - VI (published by Wiley-Interscience), or Bodansky, "The Practice of Peptide Synthesis" (published by Springer-Verlag). Hydrolysis of the ethyl ester of L-4 is accomplished according to the general conditions described for the conversion of L-2 to L-3, to provide the carboxylic acid L-5. Alternatively, the intermediate carboxylate salt of can be isolated, if desired, or a carboxylate salt of the free

20 carboxylic acid can be prepared by methods well-known to those of skill in the art.

Scheme M describes a method of preparing exemplary fibrinogen receptor templates described in Sugihara, *et al.*, EP 0529858.

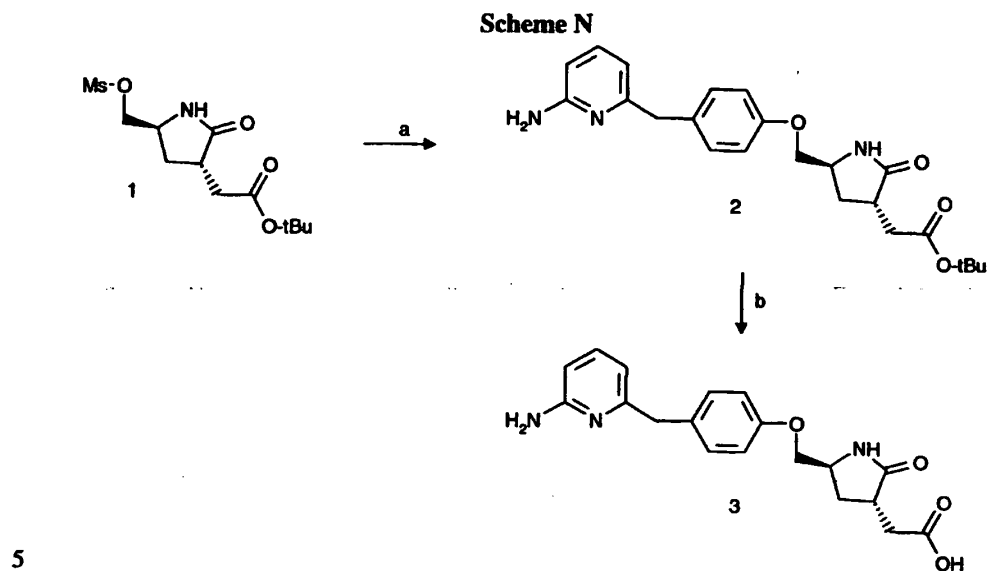


5

a) (6-amino-2-pyridinyl)acetic acid, EDC, DIEA, DMF; b) $\text{CF}_3\text{CO}_2\text{H}$, CH_2Cl_2 .

M-1, prepared as described in Sugihara, *et al.*, EP 0529858, is condensed with a
 10 suitable substituted carboxylic acid, such as (6-amino-2-pyridinyl)acetic acid, prepared by
 saponification of ethyl (6-amino-2-pyridinyl)acetate, Awaya, *et al.*, *Chem. Pharm. Bull.*
 1974, 22, 1414, to give M-2, and the tert-butyl ester is cleaved with TFA, following the
 general procedure of Sugihara, *et al.*, Example 59, to give M-3. Many additional methods
 for converting a carboxylic acid to an amide are known, and can be found in standard
 15 reference books, such as "Compendium of Organic Synthesis", Vol. I-VI (published by
 Springer-Verlag).

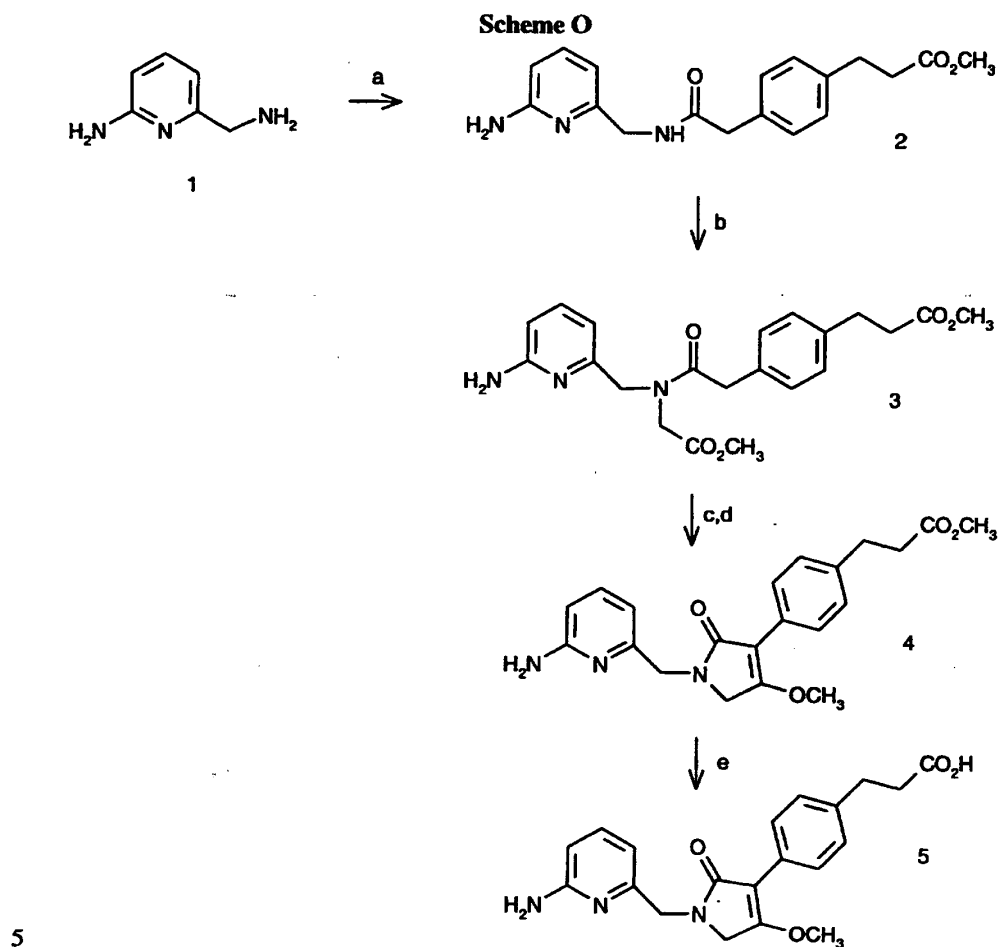
Scheme N describes a method of preparing exemplary fibrinogen receptor templates described in Himmelsbach, *et al.*, AU-A-86926/91.



a) 4-[(6-amino-2-pyridinyl)methyl]phenol, Cs_2CO_3 , DMF; b) 1N NaOH, CH_3OH .

- Compound N-1, prepared as described by Himmelsbach, *et al.*, AU-A-86926/91,
- 10 Example VI(28), is treated with a suitable substituted phenol, such as 4-[(6-amino-2-pyridinyl)methyl]phenol, prepared from the corresponding anisole, Ife, *et al.*, WO 9426715, with hydrobromic acid, following the general method of Himmelsbach *et al.*, Example 3(51), to give N-2. The tert-butyl ester in N-2 is hydrolyzed with 1N NaOH in CH_3OH to give N-3. Alternatively, the tert-butyl ester may be cleaved with TFA or HCl
- 15 in a suitable solvent such as CH_2Cl_2 .

Scheme O describes a method of preparing exemplary fibrinogen receptor templates described in Linz, *et al.*, EP 0567968.

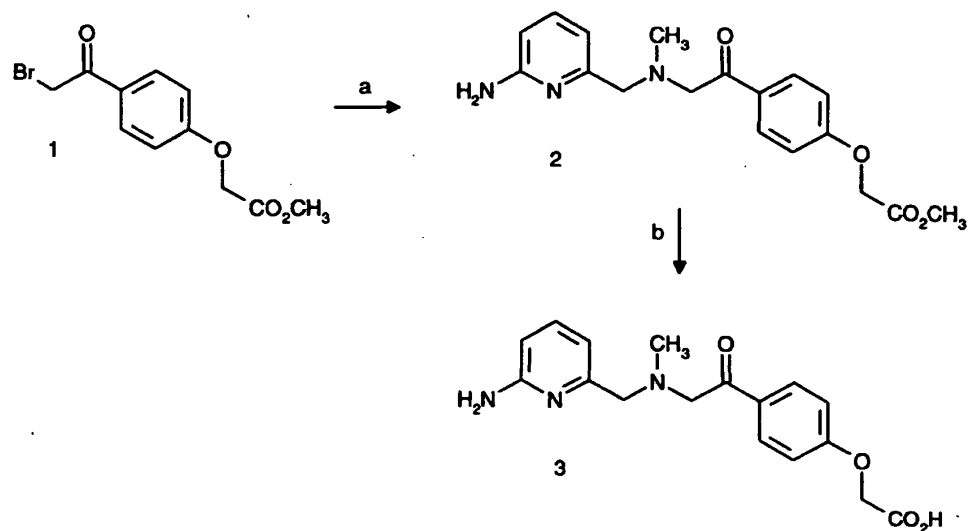


a) 6-aminomethyl-2-pyridinamine, Ph_2POCl , Et_3N , DMAP, THF; b) NaH , $\text{BrCH}_2\text{CO}_2\text{CH}_3$, DMF; c) KOtBu , CH_3I , DMF; e) LiOH , H_2O , THF.

- 10 Following the procedures of Linz, *et al.*, EP 0567968, except substituting (6-amino-2-pyridinyl)methylamine for 4-cyanoaniline, gives O-5.

Scheme P describes a method of preparing exemplary fibrinogen receptor templates described in Wayne, *et al.*, WO 94/22834.

Scheme P

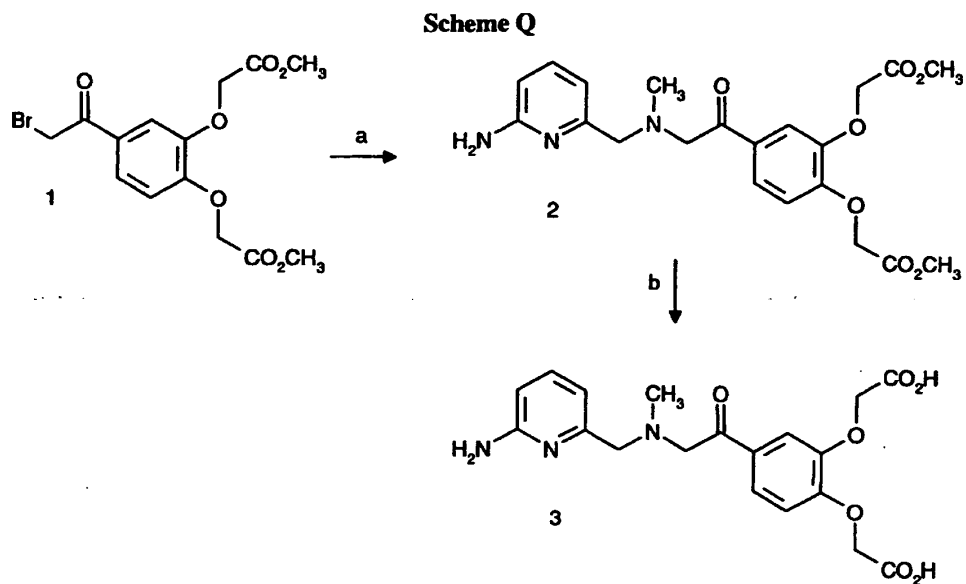


5

a) 6-(methylamino)methyl-2-pyridinamine, CH₃CN; b) 1N NaOH, CH₃OH

10 Following the procedures of Wayne, *et al.*, WO 94/22834, Example 1-2, except substituting 6-(methylamino)methyl-2-pyridinamine for 1-(4-pyridyl)piperazine gives **P-3**.

Scheme Q describes a method of preparing exemplary fibrinogen receptor templates described in Wayne, *et al.*, WO 94/22834.

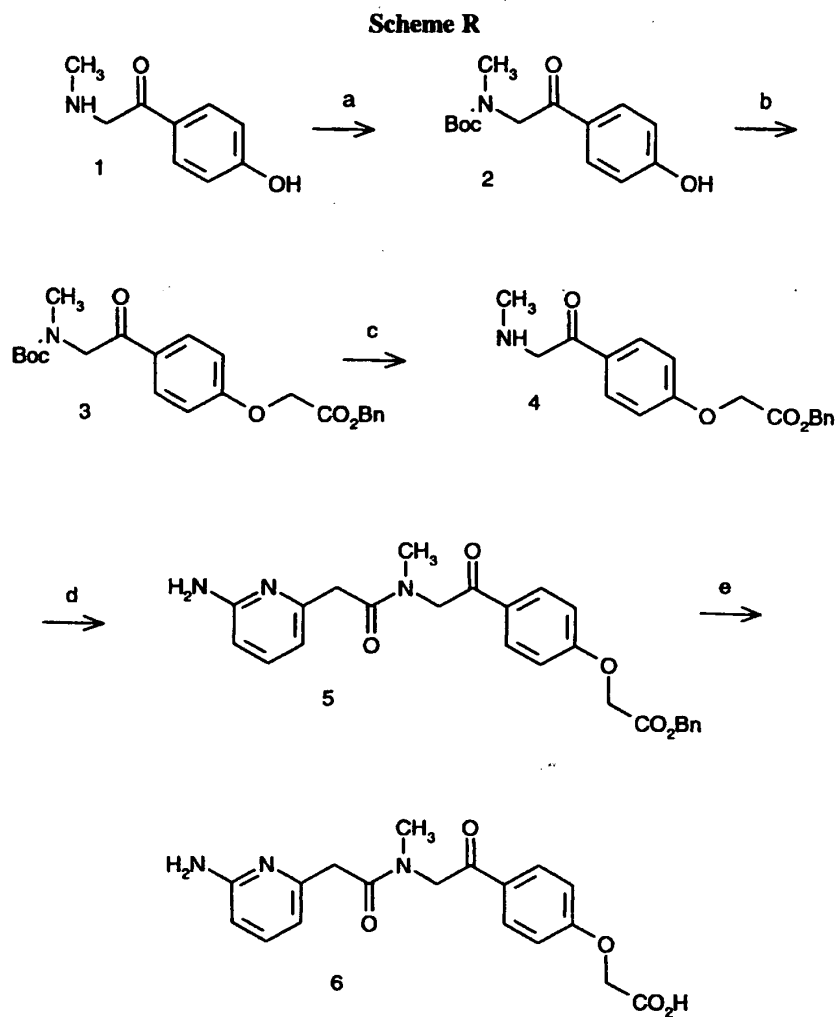


5

a) 6-(methylamino)methyl-2-pyridinamine, CH₃CN; b) 1N NaOH, CH₃OH

10 Following the procedures of Wayne, *et al.*, WO 94/22834, Example 3-4, except substituting 6-(methylamino)methyl-2-pyridinamine for 1-(4-pyridyl)piperazine gives Q-3.

Scheme R describes a method of preparing exemplary fibrinogen receptor templates described in Alig, *et al.*, EP 0381033.



a) (Boc)₂O, NaOH, dioxane, H₂O; b) BrCH₂CO₂Bn, K₂CO₃, acetone; c) 4M HCl, dioxane; d) (6-amino-2-pyridinyl)acetic acid, EDC, DIEA, DMF; e) 1N NaOH, CH₃OH.

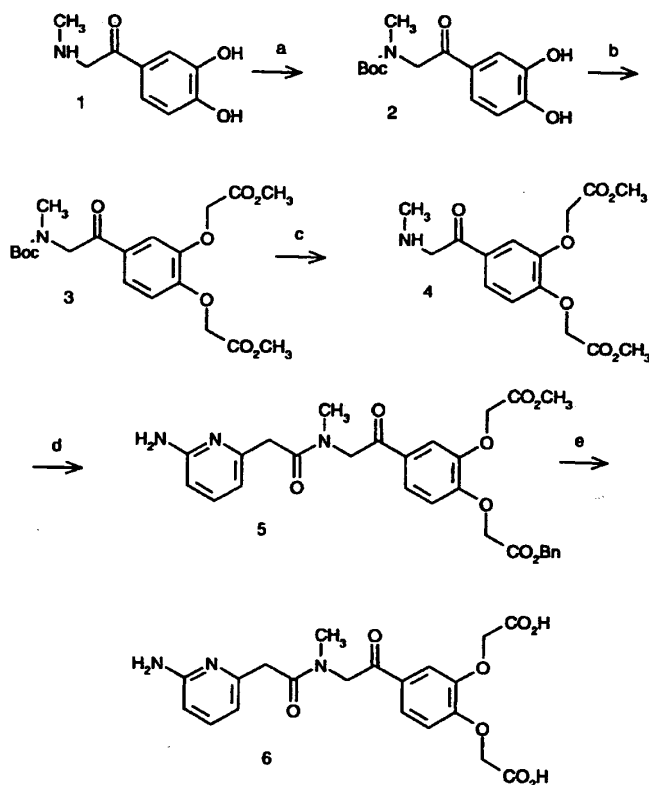
- 10 **R-1** is treated with di-tert-butyl dicarbonate and sodium hydroxide in aqueous dioxane to afford **R-2**, which is alkylated on the phenolic oxygen with benzyl bromoacetate and potassium carbonate in acetone to give **R-3**. The Boc group in **R-3** is

removed with hydrogen chloride in dioxane, and the resulting **R-4** is acylated on nitrogen with (6-amino-2-pyridinyl)acetic acid, prepared by saponification of ethyl (6-amino-2-pyridinyl)acetate, Awaya, *et al.*, *Chem. Pharm. Bull.* **1974**, *22*, 1414, EDC and DIEA in DMF to give **R-5**. The benzyl ester in **R-5** is saponified to give **R-6**. Alternatively, the benzyl ester may be cleaved by treatment with H₂ and a suitable catalyst, such as Pd/C, in a suitable solvent, such as CH₃OH, EtOH, or AcOH.

Scheme S describes a method of preparing exemplary fibrinogen receptor templates described in Alig, *et al.*, EP 0381033.

10

Scheme S



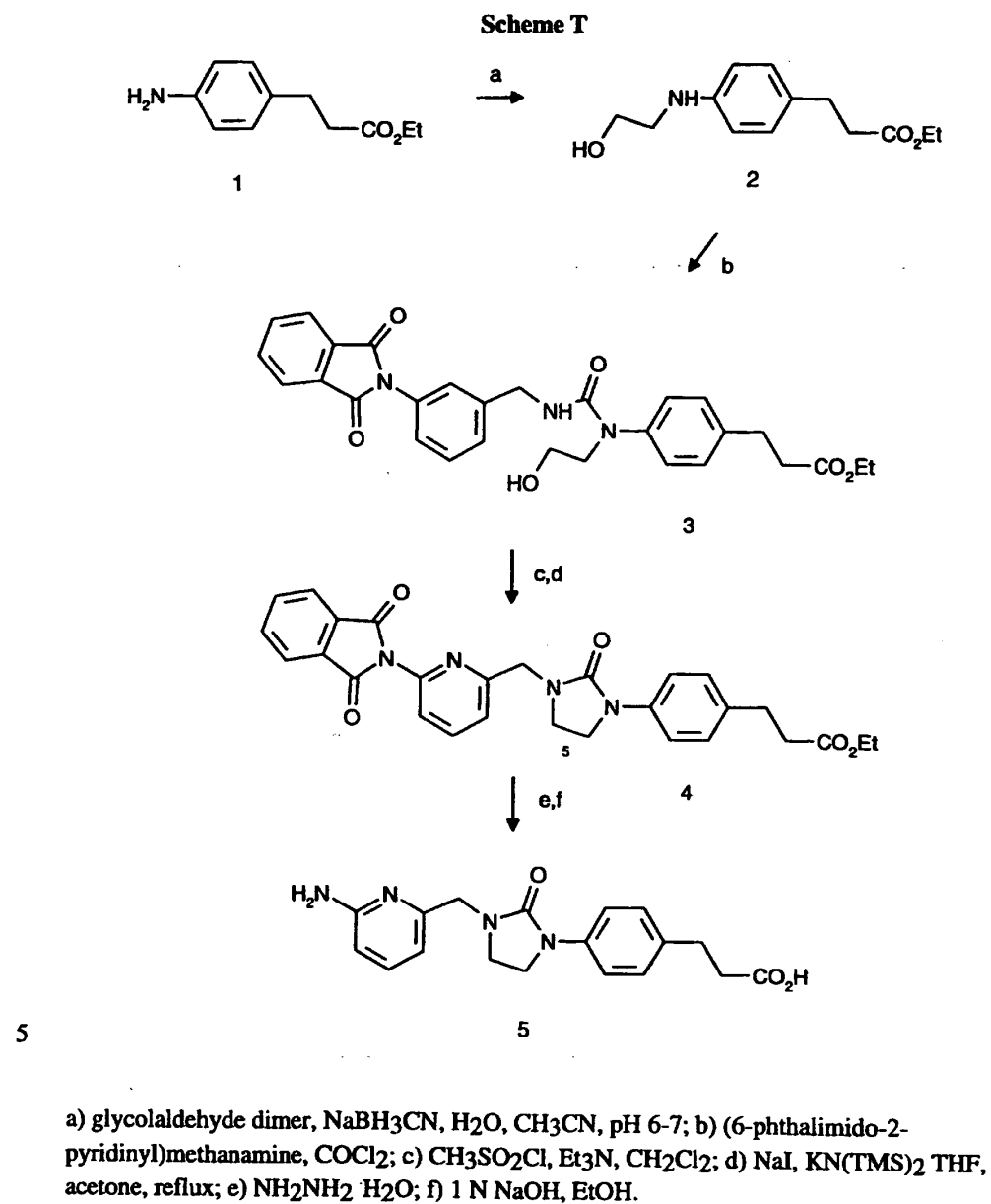
a) (Boc)₂O, NaOH, dioxane, H₂O; b) BrCH₂CO₂CH₃, K₂CO₃, acetone; c) 4M HCl, dioxane; d) (6-amino-2-pyridinyl)acetic acid, EDC, DIEA, DMF; e) 1N NaOH, CH₃OH.

15

S-1 is treated with di-tert-butyl dicarbonate and sodium hydroxide in aqueous dioxane to afford **S-2**, which is alkylated on the phenolic oxygens with methyl

- bromoacetate and potassium carbonate in acetone to give S-3. The Boc group in S-3 is removed with hydrogen chloride in dioxane, and the resulting S-4 is acylated on nitrogen with (6-amino-2-pyridinyl)acetic acid, prepared by saponification of ethyl (6-amino-2-pyridinyl)acetate, Awaya, *et al.*, *Chem. Pharm. Bull.* **1974**, 22, 1414, EDC and DIEA in
- 5 DMF to give S-5. The methyl esters in R-5 are cleaved by treatment with 1M NaOH in CH₃OH to give S-6.

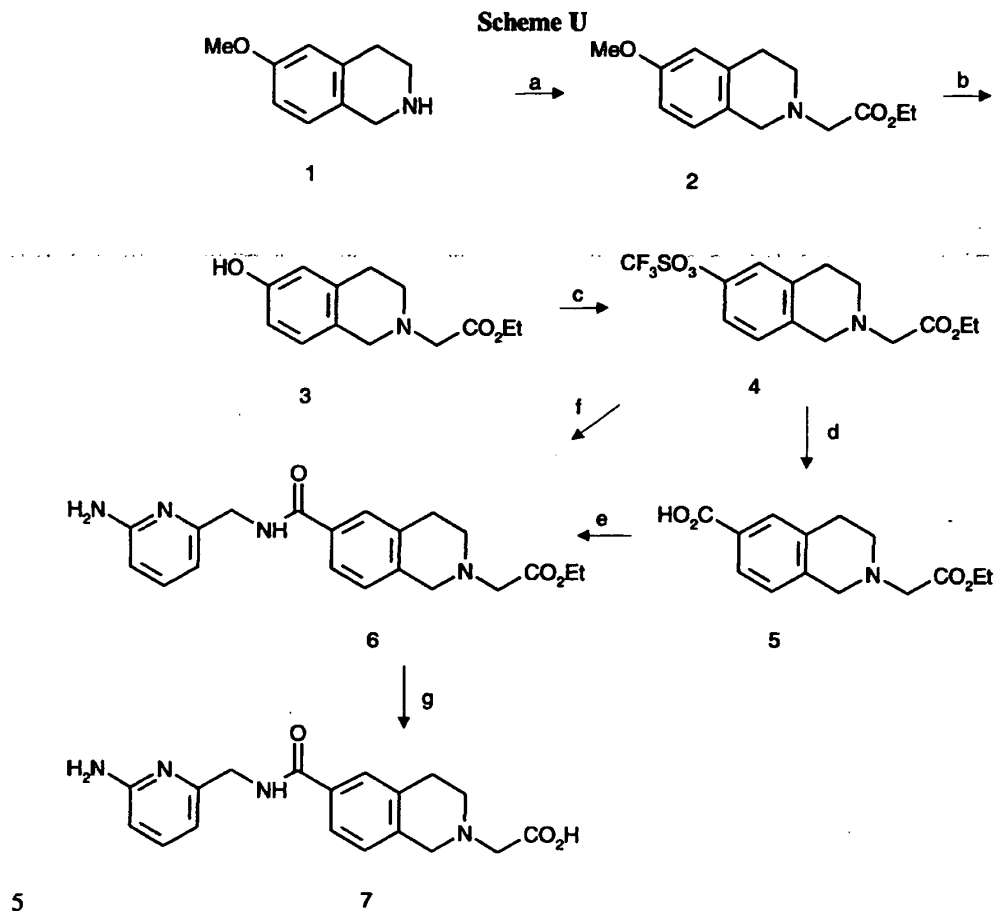
Scheme T describes a method of preparing exemplary fibrinogen receptor templates described in Himmelsbach, *et al.*, EP 0587134.



10

Scheme T provides a method for the preparation of 2-oxo-imidazolidine compounds, e.g., **T-5**, wherein reductive amination of an amine, for example **T-1**, with glycolaldehyde dimer and sodium cyanoborohydride, gives a secondary amine, such as **T-2**. A primary amine, as exemplified by (6-phthalimido-2-pyridinyl)methanamine, is
5 treated with phosgene to give an isocyanate, which is allowed to react, without isolation, with the secondary hydroxyethylamine to give a hydroxyethylurea, as exemplified by compound **T-3**. The hydroxyl group is converted into a leaving group, such as a methanesulfonate or iodide, and is allowed to cyclize to a 2-oxo-imidazolidine, **T-4**,
employing methods known in the art, Himmelsbach, *et al.*, EP 0587134, such as treating
10 the hydroxyethylurea **4** with trifluorosulfonyl chloride and Et₃N, followed by NaI and then potassium bis(trimethylsilyl)azide, as described in Himmelsbach, *et al.*, EP 0587134, Example III. Treatment of **T-4** with hydrazine and saponification of the ester give **T-5**.

Scheme U provides a method for the preparation of 1,2,3,4-tetrahydroisoquinoline compounds as exemplary fibrinogen receptor antagonists, as described in M. J. Fisher *et al.*, EP 0635492.



- a) $\text{ClCH}_2\text{CO}_2\text{Et}$, Et_3N , DMF; b) BBr_3 , CH_2Cl_2 ; c) $(\text{CF}_3\text{SO}_2)_2\text{O}$, pyridine; d) CO, $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O ; e) (6-amino-2-pyridinyl)methanamine, EDC, HOBT, DIEA, DMF; f) (6-amino-2-pyridinyl)methanamine, CO, $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O ; g) 1N NaOH, EtOH.
- 10

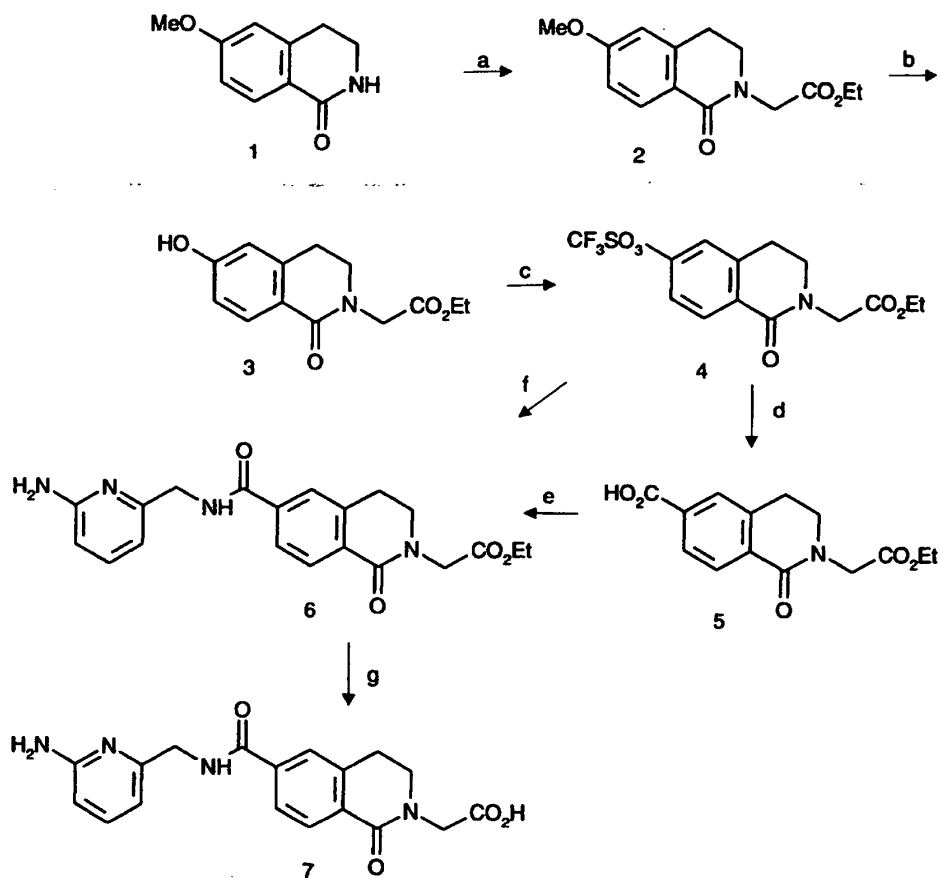
Accordingly, a 6-methoxy-3,4-dihydroisoquinoline, such as compound U-1 is prepared by the method described by D. J. Sall and G. L. Grunewald, *J. Med. Chem.* **1987**, *30*, 2208-2216. The isoquinoline is treated with a haloacetic acid ester in the

presence of a tertiary amine to afford the 2-acetic acid ester, as exemplified by compound U-2. The 6-methoxy compound is converted into the corresponding 6-hydroxy compound by methods known in the art, for example with BBr₃, which is converted into the triflate with trifluorosulfonic acid anhydride. Palladium-catalyzed carbonylation affords the 6-
5 carboxy compound, such as compound U-5, which is then condensed with an amine, as exemplified by (6-amino-2-pyridinyl)methanamine, employing a standard amide bond forming reagent to give the desired amide, such as compound U-6. Saponification affords the title compound of Example W, U-7. Alternatively, the palladium-catalyzed
10 carbonylation reaction with the triflate, exemplified by compound U-4, may be trapped with (6-amino-2-pyridinyl)methanamine to provide, after saponification, the compound of Example W, U-7.

Scheme V provides a method for the preparation of 3,4-dihydroisoquinolin-1-one compounds as exemplary fibrinogen receptor antagonists, as described M. J. Fisher *et al.*, EP 0635492.

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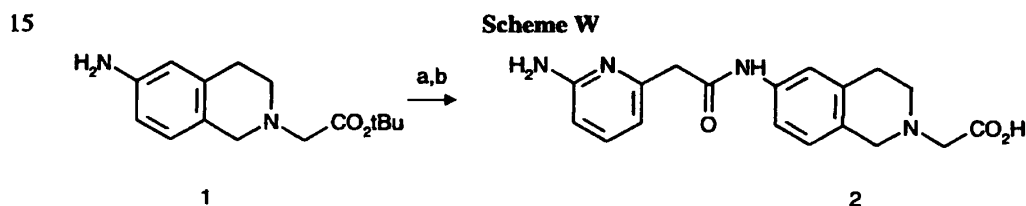
Scheme V



- a) 1. $\text{LiN}(\text{TMS})_2$, 2. $\text{ClCH}_2\text{CO}_2\text{Et}$, DMF; b) BBr_3 , CH_2Cl_2 ; c) $(\text{CF}_3\text{SO}_2)_2\text{O}$, pyridine; d) CO , $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O ; e) N-(2-pyridinyl)ethylenediamine, EDC, HOBt, DIEA, DMF; f) (6-amino-2-pyridinyl)methanamine, CO , $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O ; g) 1N NaOH, EtOH.

Accordingly, the 1-oxo compound V-1, prepared by the method described by D. J. Sall and G. L. Grunewald, *J. Med. Chem.* **1987**, *30*, 2208-2216, is treated with a base, such as LiN(TMS)₂, and a haloacetic acid ester to give a 2-acetic acid ester, as exemplified by compound V-2. The 1-oxo compound is then employed in the analogous series of reactions deployed in Scheme U, substituting the corresponding 1-oxo analog, as shown in Scheme U, to provide the title compound of Example X, V-7. As in Scheme U, alternatively, the palladium-catalyzed carbonylation reaction with the triflate, exemplified by compound V-4, may be trapped with an amine, such as (6-amino-2-pyridinyl)methanamine, provides, after saponification, the amide exemplified by the title compound of Example X, V-7.

Scheme W provides a method for the preparation of 6-acylaminotetralin compounds as exemplary fibrinogen receptor antagonists, as described M. J. Fisher *et al.*, EP 0635492.



20 a) (6-amino-2-pyridinyl)acetic acid, EDC, HOBt, DIEA, DMF; b) TFA, CH₂Cl₂.

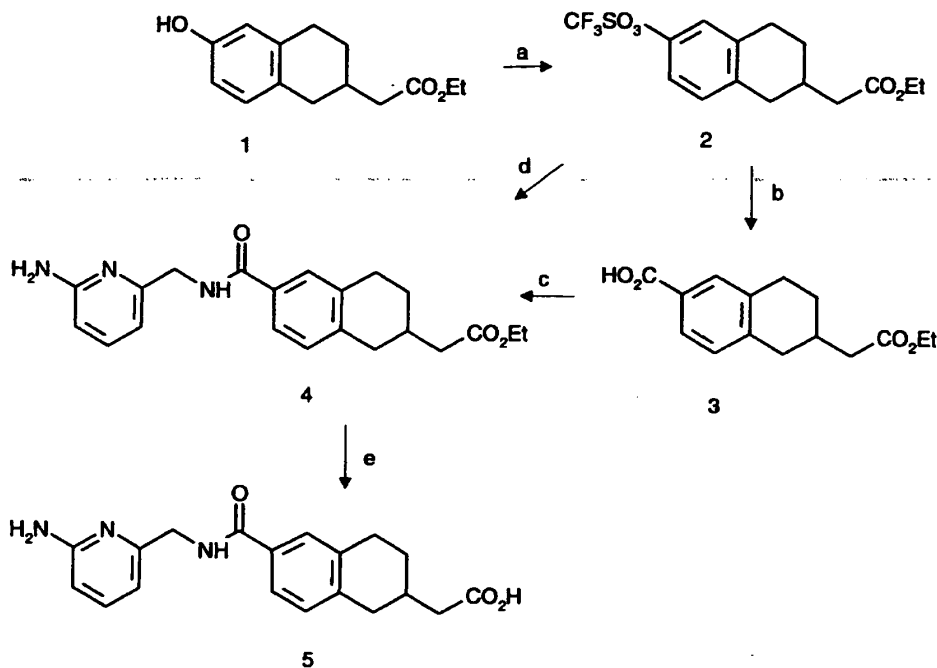
Accordingly, a 6-amino-2-tert-butoxycarbonyl-tetral-1-one, exemplified by compound W-1, which is prepared according to the methods described in M. J. Fisher *et al.*, EP 0635492, is condensed with an activated derivative of a carboxylic acid obtained (6-amino-2-pyridinyl)acetic acid to provide, after deesterification, the amide exemplified by the title compound of Example Y, W-2.

25

Scheme X provides a method for the preparation of 6-aminoacyltetralin compounds as exemplary fibrinogen receptor antagonists, as described M. J. Fisher *et al.*, EP 0635492.

5

Scheme X

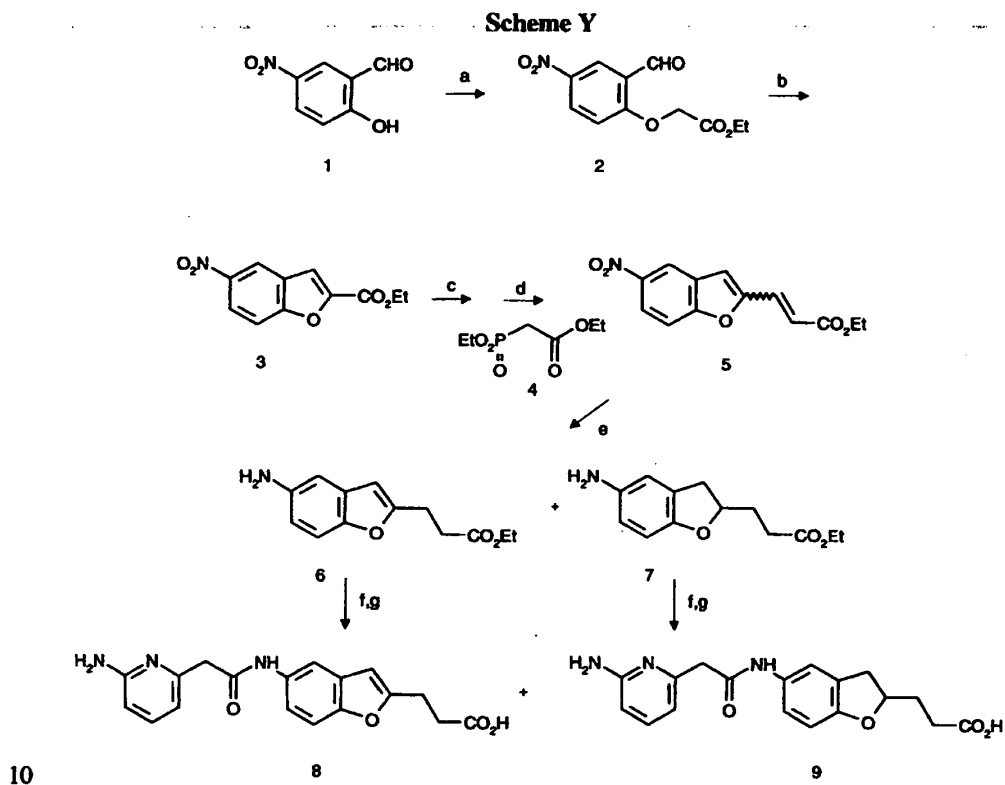


- a) $(\text{CF}_3\text{SO}_2)\text{O}$, pyridine; b) CO , $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O ; c) (6-amino-2-pyridinyl)methanamine, EDC, HOBt, DIEA, DMF; d) (6-amino-2-pyridinyl)methanamine, CO , $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O ; e) 1N NaOH, EtOH.

Accordingly, an ethoxycarbonylmethyl-6-hydroxy-tetral-1-one, exemplified by compound X-1, which is prepared according to the methods described in M. J. Fisher *et al.*, EP 0635492, is treated with triflic anhydride to provide the triflate, as exemplified by compound X-2, which is employed in a palladium-catalyzed carbonylation reaction to afford a carboxylic acid, such as compound X-3, which is then condensed with an amine such as (6-amino-2-pyridinyl)methanamine to provide, after deesterification, the 6-

aminoacyl compound exemplified by Example Z, X-5. Alternatively, the palladium-catalyzed carbonylation reaction with the triflate exemplified by compound X-2, may be trapped with (6-amino-2-pyridinyl)methanamine to provide, after saponification, the corresponding 6-aminoacyl compound X-5.

- 5 Scheme Y provides a method for the preparation of 5-acylaminobenzofuran and 5-acylaminodihydrobenzofuran compounds as exemplary fibrinogen receptor antagonists, as described in M. L. Denney, *et al.*, EP 0655439.



a) $\text{BrCH}_2\text{CO}_2\text{Et}$, K_2CO_3 , NaI , THF; b) 1. DBU, EtOH, 2. HCl , EtOH; c) DiBAL, -78°C , THF; d) NaH , THF; e) H_2 , 10% Pd/C , EtOH; f) (6-amino-2-pyridinyl)acetic acid, EDC, HOBT, Et_3N , DMF; g) 1N NaOH , CH_3OH .

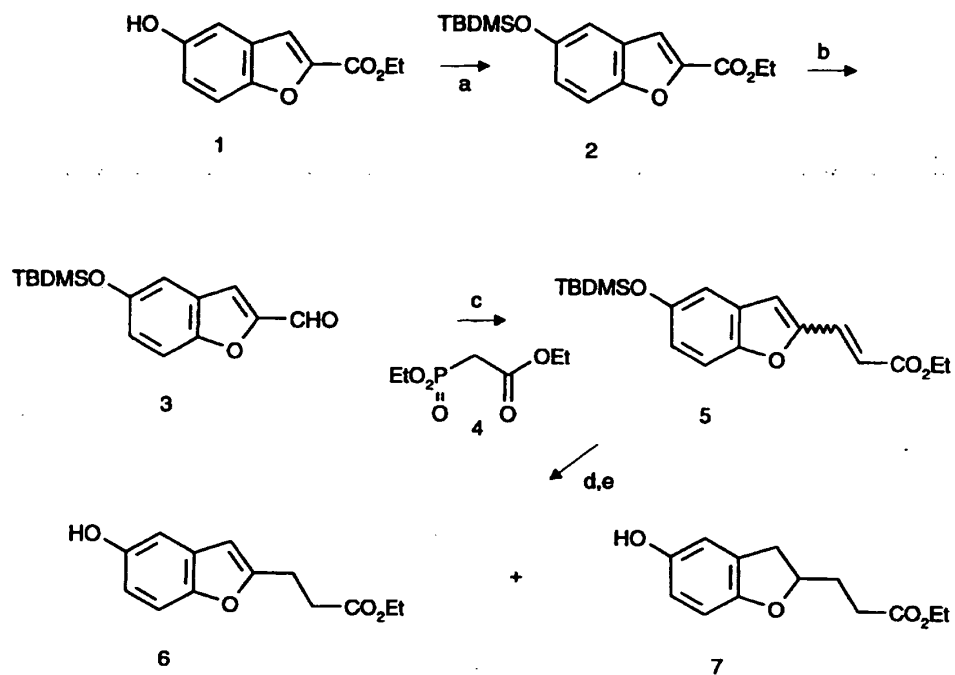
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Accordingly, a 5-nitrosalicylaldehyde, exemplified by compound Y-1, is treated with a haloacetic acid ester to give the phenoxyacetic acid ester, exemplified by compound Y-2. A 2-alkoxycarbonylfuran, exemplified by compound Y-3, is obtained by treating the aldehyde with base, for example with DBU. The 2-alkoxycarbonyl group is
5 reduced to the aldehyde, for example with DiBAL. Wittig reaction affords the 2-acrylate ester, exemplified by compound Y-5, which is reduced to the benzofuran-2-propionic acid ester, exemplified by compound Y-6 and the dihydrobenzofuran-2-propionic acid ester, exemplified by compound Y-7. The amine Y-6 is then condensed with an activated derivative of a carboxylic acid, such as (6-amino-2-pyridinyl)acetic acid, to provide, after
10 deesterification, the amide exemplified by the title compound of Example AA, Y-8. Alternatively, the amine Y-7 is condensed with an activated derivative of a carboxylic acid, such as (6-amino-2-pyridinyl)acetic acid, to provide, after deesterification, the amide Y-9.

Schemes Z-1, Z-2 and Z-3 provide a method for the preparation of 5-aminoacylbenzofuran and 5-aminoacyldihydrobenzofuran compounds as exemplary fibrinogen receptor antagonists, as described in M. L. Denney, *et al.*, EP 0655439.

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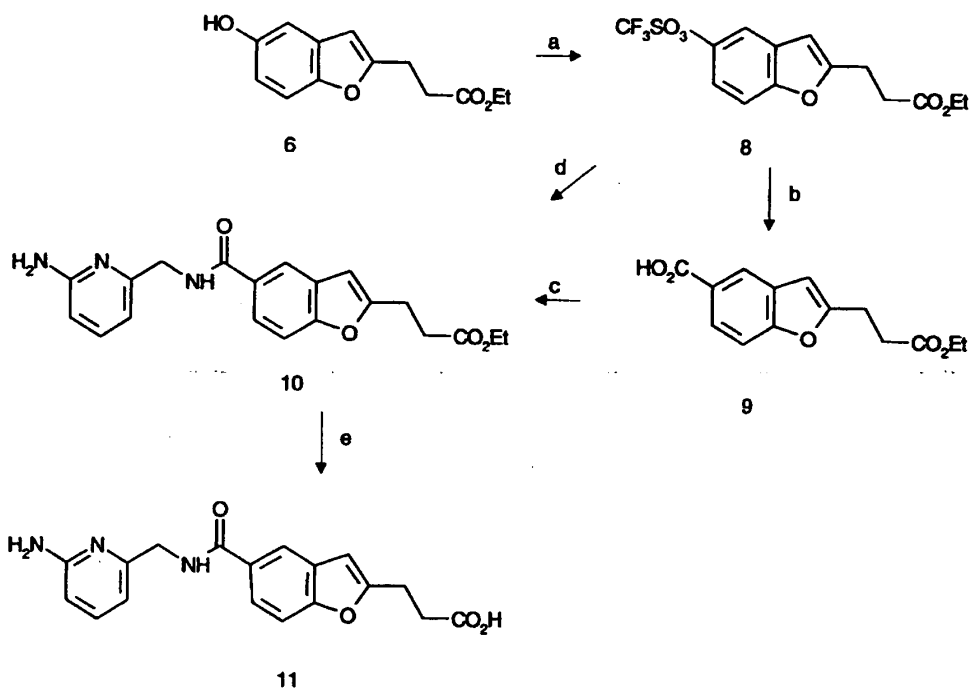
Scheme Z-1



a) TBDMS-Cl, imidazole, THF; b) DiBAL, -78°C, THF; c) NaH, THF; d) H₂, 5% Pd/C, EtOH; e) Et₄NF, THF.

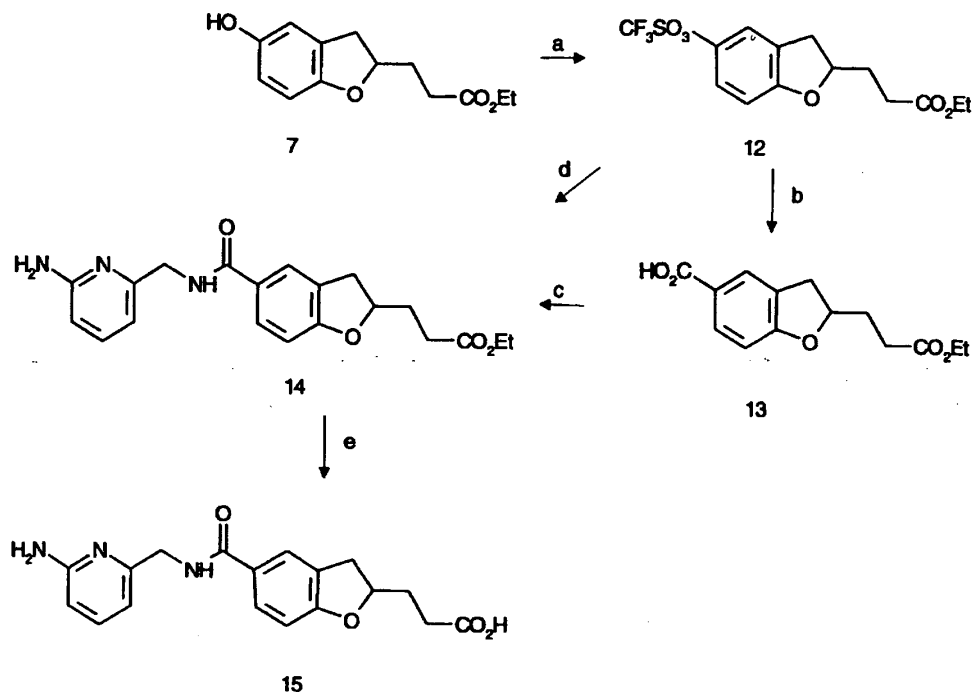
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Scheme Z-2



- 5 a) $(\text{CF}_3\text{SO}_2)_2\text{O}$, pyridine; b) CO, $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O ; c) (6-amino-2-pyridinyl)methanamine, EDC, HOBT, DIEA, DMF; d) (6-amino-2-pyridinyl)methanamine, CO, $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O e) 1N NaOH, EtOH.

Scheme Z-3

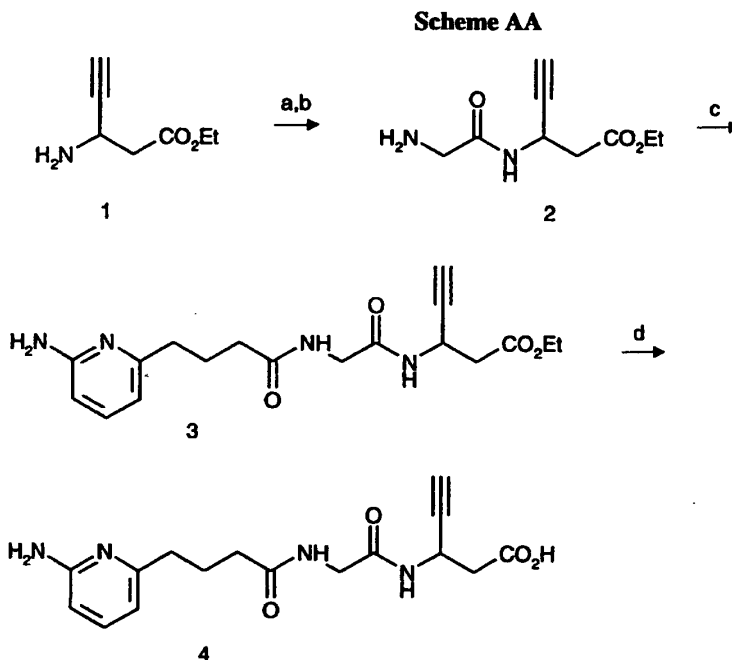


- 5 a) $(\text{CF}_3\text{SO}_2)_2\text{O}$, pyridine; b) CO, $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O ; c) (6-amino-2-pyridinyl)methanamine, EDC, HOBT, DIEA, DMF; d) (6-amino-2-pyridinyl)methanamine, CO, $\text{Pd}(\text{OAc})_2$, PPh_3 , DIEA, NMP, NH_4HCO_3 , H_2O ; e) 1N NaOH, EtOH.

- 10 Accordingly, a 5-hydroxybenzofuran-2-carboxylic acid ester, such as compound Z-1-1, prepared in the manner of M. L. Denney, *et al.*, EP 0655439, is treated with TBDMS-Cl to provide the TBDMS derivative of the ester, Z-1-2. The ester is reduced to an aldehyde, such as compound Z-1-3. Wittig reaction affords an acrylic acid ester, as exemplified by compound Z-1-5. Catalytic reduction affords a benzofuran-2-acetic acid ester and a dihydrobenzofuran-2-acetic acid ester. Cleavage of the silyl ether group of each ester, by methods known to the art, affords a benzofuran-2-acetic acid ester, as exemplified by compound Z-1-6 and a dihydrobenzofuran-2-acetic acid ester as exemplified by compound Z-1-7.
- 15

As shown in Schemes Z-2 and Z-3, each phenol may be converted to a carboxylic acid via palladium-catalyzed carbonylation, such as compound **Z-2-9** or **Z-3-13**, which are then condensed with an amine, such as (6-amino-2-pyridinyl)methanamine, to provide, after deesterification, the amide of the title compound of Example CC (**Z-2-11**) or DD (**Z-3-15**). Alternatively, the palladium-catalyzed carbonylation reaction with the triflates exemplified by compounds **Z-2-8**, or **Z-3-12**, may be trapped with (6-amino-2-pyridinyl)methanamine to provide, after deesterification, the corresponding 6-aminoacyl compounds, Example CC (**Z-2-11**) or DD (**Z-3-15**).

Scheme AA describes a method of preparing a further exemplary fibrinogen receptor template.



- 15 a) Boc-Gly, EDC, HOBT, DIEA, CH₃CN; b) TFA, CH₂Cl₂; c) 4-(6-amino-2-pyridinyl)butanoic acid, EDC, HOBT, DIEA, DMF; d) 1N LiOH, THF, CH₃CN.

The preparation of the intermediate **AA-2** begins with the coupling of the known ethyl 3-amino-4-pentynoate (WO 93/07867) with commercially available tert-

butoxycarbonylglycine (Boc-Gly) under standard peptide bond forming conditions. The product of this reaction is deprotected to AA-2 under acidic conditions which are known to effect removal of a Boc protecting group. The two intermediates AA-2 and 4-(6-amino-2-pyridinyl)butanoic acid are coupled under standard peptide coupling conditions to give AA-3, which is hydrolyzed to AA-4 with lithium hydroxide in aqueous THF and CH₃CN.

Acid addition salts of the compounds are prepared in a standard manner in a suitable solvent from the parent compound and an excess of an acid, such as hydrochloric, hydrobromic, hydrofluoric, sulfuric, phosphoric, acetic, trifluoroacetic, maleic, succinic or methanesulfonic. Certain of the compounds form inner salts or zwitterions which may be acceptable. Cationic salts are prepared by treating the parent compound with an excess of an alkaline reagent, such as a hydroxide, carbonate or alkoxide, containing the appropriate cation; or with an appropriate organic amine. Cations such as Li⁺, Na⁺, K⁺, Ca⁺⁺, Mg⁺⁺ and NH₄⁺ are specific examples of cations present in pharmaceutically acceptable salts.

This invention also provides a pharmaceutical composition which comprises a compound according to formula (I) and a pharmaceutically acceptable carrier. Accordingly, the compounds of formula (I) may be used in the manufacture of a medicament. Pharmaceutical compositions of the compounds of formula (I) prepared as hereinbefore described may be formulated as solutions or lyophilized powders for parenteral administration. Powders may be reconstituted by addition of a suitable diluent or other pharmaceutically acceptable carrier prior to use. The liquid formulation may be a buffered, isotonic, aqueous solution. Examples of suitable diluents are normal isotonic saline solution, standard 5% dextrose in water or buffered sodium or ammonium acetate solution. Such formulation is especially suitable for parenteral administration, but may also be used for oral administration or contained in a metered dose inhaler or nebulizer for insufflation. It may be desirable to add excipients such as polyvinylpyrrolidone, gelatin, hydroxy cellulose, acacia, polyethylene glycol, mannitol, sodium chloride or sodium citrate.

Alternately, these compounds may be encapsulated, tableted or prepared in a emulsion or syrup for oral administration. Pharmaceutically acceptable solid or liquid carriers may be added to enhance or stabilize the composition, or to facilitate preparation of the composition. Solid carriers include starch, lactose, calcium sulfate dihydrate, terra alba, magnesium stearate or stearic acid, talc, pectin, acacia, agar or gelatin. Liquid carriers include syrup, peanut oil, olive oil, saline and water. The carrier may also include a sustained release material such as glyceryl monostearate or glyceryl distearate, alone or with a wax. The amount of solid carrier varies but, preferably, will be between

about 20 mg to about 1 g per dosage unit. The pharmaceutical preparations are made following the conventional techniques of pharmacy involving milling, mixing, granulating, and compressing, when necessary, for tablet forms; or milling, mixing and filling for hard gelatin capsule forms. When a liquid carrier is used, the preparation will be in the form of a syrup, elixir, emulsion or an aqueous or non-aqueous suspension. Such a liquid formulation may be administered directly p.o. or filled into a soft gelatin capsule.

For rectal administration, the compounds of this invention may also be combined with excipients such as cocoa butter, glycerin, gelatin or polyethylene glycols and molded into a suppository.

The compounds described herein are antagonists of the vitronectin receptor, and are useful for treating diseases wherein the underlying pathology is attributable to ligand or cell which interacts with the vitronectin receptor. For instance, these compounds are useful for the treatment of diseases wherein loss of the bone matrix creates pathology. Thus, the instant compounds are useful for the treatment of osteoporosis, hyperparathyroidism, Paget's disease, hypercalcemia of malignancy, osteolytic lesions produced by bone metastasis, bone loss due to immobilization or sex hormone deficiency. The compounds of this invention are also believed to have utility as antitumor, antiinflammatory, anti-angiogenic and anti-metastatic agents, and be useful in the treatment of cancer, atherosclerosis and restenosis.

The peptide is administered either orally or parenterally to the patient, in a manner such that the concentration of drug is sufficient to inhibit bone resorption, or other such indication. The pharmaceutical composition containing the peptide is administered at an oral dose of between about 0.1 to about 50 mg/kg in a manner consistent with the condition of the patient. Preferably the oral dose would be about 0.5 to about 20 mg/kg. For acute therapy, parenteral administration is preferred. An intravenous infusion of the peptide in 5% dextrose in water or normal saline, or a similar formulation with suitable excipients, is most effective, although an intramuscular bolus injection is also useful. Typically, the parenteral dose will be about 0.01 to about 100 mg/kg; preferably between 0.1 and 20 mg/kg. The compounds are administered one to four times daily at a level to achieve a total daily dose of about 0.4 to about 400 mg/kg/day. The precise level and method by which the compounds are administered is readily determined by one routinely skilled in the art by comparing the blood level of the agent to the concentration required to have a therapeutic effect.

The compounds may be tested in one of several biological assays to determine the concentration of compound which is required to have a given pharmacological effect.

INHIBITION OF VITRONECTIN BINDING

Solid-Phase [³H]-SK&F-107260 Binding to $\alpha_v\beta_3$: Human placenta or human platelet $\alpha_v\beta_3$ (0.1-0.3 mg/mL) in buffer T (containing 2 mM CaCl₂ and 1% octylglucoside) was diluted with buffer T containing 1 mM CaCl₂, 1 mM MnCl₂, 1 mM MgCl₂ (buffer A) and 0.05% NaN₃, and then immediately added to 96-well ELISA plates (Corning, New York, NY) at 0.1 mL per well. 0.1 - 0.2 μ g of $\alpha_v\beta_3$ was added per well. The plates were incubated overnight at 4°C. At the time of the experiment, the wells were washed once with buffer A and were incubated with 0.1 mL of 3.5% bovine serum albumin in the same buffer for 1 hr at room temperature. Following incubation the wells were aspirated completely and washed twice with 0.2 mL buffer A.

Compounds were dissolved in 100% DMSO to give a 2 mM stock solution, which was diluted with binding buffer (15 mM Tris-HCl (pH 7.4), 100 mM NaCl, 1 mM CaCl₂, 1 mM MnCl₂, 1 mM MgCl₂) to a final compound concentration of 100 μ M. This solution is then diluted to the required final compound concentration. Various concentrations of unlabeled antagonists (0.001 - 100 μ M) were added to the wells in triplicates, followed by the addition of 5.0 nM of [³H]-SK&F-107260 (65 - 86 Ci/mmol).

The plates were incubated for 1 hr at room temperature. Following incubation the wells were aspirated completely and washed once with 0.2 mL of ice cold buffer A in a well-to-well fashion. The receptors were solubilized with 0.1 mL of 1% SDS and the bound [³H]-SK&F-107260 was determined by liquid scintillation counting with the addition of 3 mL Ready Safe in a Beckman LS Liquid Scintillation Counter, with 40% efficiency. Nonspecific binding of [³H]-SK&F-107260 was determined in the presence of 2 μ M SK&F-107260 and was consistently less than 1% of total radioligand input. The IC₅₀ (concentration of the antagonist to inhibit 50% binding of [³H]-SK&F-107260) was determined by a nonlinear, least squares curve-fitting routine, which was modified from the LUNDON-2 program. The K_i (dissociation constant of the antagonist) was calculated according to the equation: $K_i = IC_{50}/(1 + L/K_d)$, where L and K_d were the concentration and the dissociation constant of [³H]-SK&F-107260, respectively.

Compounds of the present invention inhibit vitronectin binding to SK&F 107260 in the concentration range of 0.01 to 25 micromolar. Preferred compounds inhibit vitronectin binding at a concentration of less than 1 micromolar.

Compounds of this invention are also tested for *in vitro* and *in vivo* bone resorption in assays standard in the art for evaluating inhibition of bone formation, such as the pit formation assay disclosed in EP 528 587, which may also be performed using human osteoclasts in place of rat osteoclasts, and the ovariectomized rat model, described by Wronski *et al.*, *Cells and Materials* 1991, Sup. 1, 69-74.

PARATHYROIDECTOMIZED RAT MODEL

- Each experimental group consists of 5-6 male Sprague-Dawley rats. The rats are parathyroidectomized (by the vendor, Taconic Farms) 7 days prior to use. Twenty four hours prior to use, circulating ionized calcium levels are measured in whole blood immediately after it has been withdrawn by tail venipuncture into heparinized tubes. Rats are included if ionized Ca level (measured with a Ciba-Corning model 634 calcium pH analyzer) is ≥ 1.2 mM/L. The rats are then put on a diet of calcium-free chow and deionized water. At the start of the experiment the rats weigh approximately 100g. Baseline Ca levels are measured and the rats are administered control vehicle (saline) or compound (dissolved in saline) as a single intravenous (tail vein) bolus injection followed immediately by a single subcutaneous injection of either human parathyroid hormone 1-34 peptide (hPTH1-34, dose 0.2mg/kg in saline/0.1% bovine serum albumen, Bachem, Ca) or the PTH vehicle. The calcemic response to PTH (and any effect of compound on this response) is measured 2h after compound/PTH administration.

15 RAT ULNA DRIFT MODEL

- Each experimental group consists of 8-10 male Sprague-Dawley or Wistar rats of approximately 30-40g body weight at the start of the experiment. The agent being tested is administered by an appropriate route as single or multiple daily doses for a period of seven days. Prior to administration of the first dose, the rats are given a single dose of a fluorescent marker (tetracycline 25mg/kg, or calcein 10mg/kg) that labels the position of bone forming surfaces at that point in time. After dosing of compound has been completed, the rats are killed and both forelimbs are removed at the elbow, the foot is removed at the ankle and the skin removed. The sample is frozen and mounted vertically on a microtome chuck. Cross sections of the midshaft region of the ulna are cut in the cryostat. The rate of bone resorption is measured morphometrically in the medial-dorsal portion of the cortical bone. The measurement is done as follows: the amount of bone resorbed at the periosteal surface is equal to the distance by which the periosteal surface has advanced towards the fluorescent label which had been incorporated at the endosteal bone formation surface on day zero; this distance is calculated by subtracting the width of bone between the label and the periosteal surface on day 7 from the width on day zero; the resorption rate in microns per day is calculated by dividing the result by 7.

HUMAN OSTEOCLAST RESORPTION ASSAY ("PIT ASSAY")

- Aliquots of osteoclastoma-derived cell suspensions are removed from liquid nitrogen storage, warmed rapidly at 37°C and washed x1 in RPMI-1640 medium by centrifugation (1000rpm, 5 mins at 4°C).

- Aspirate the medium and replace it with murine anti-HLA-DR antibody, diluted 1:3 in RPMI-1640 medium. Incubate for 30 mins on ice and mix the cell suspension frequently.
- The cells are washed x2 with cold RPMI-1640 by centrifugation (1000rpm, 5 mins at 4°C) and the cells are transferred to a sterile 15 ml centrifuge tube. The number of
- 5 mononuclear cells are enumerated in an improved Neubauer counting chamber.
- Sufficient magnetic beads (5 / mononuclear cell), coated with goat anti-mouse IgG, are removed from their stock bottle and placed into 5 ml of fresh medium (this washes away the toxic azide preservative). The medium is removed by immobilizing the beads on a magnet and is replaced with fresh medium.
- 10 • The beads are mixed with the cells and the suspension is incubated for 30 mins on ice. The suspension is mixed frequently.
- The bead-coated cells are immobilized on a magnet and the remaining cells (osteoclast-rich fraction) are decanted into a sterile 50 ml centrifuge tube.
- Fresh medium is added to the bead-coated cells to dislodge any trapped osteoclasts. This
- 15 wash process is repeated x10. The bead-coated cells are discarded.
- The osteoclasts are enumerated in a counting chamber, using a large-bore disposable plastic pasteur to charge the chamber with the sample.
- The cells are pelleted by centrifugation and the density of osteoclasts adjusted to
- 20 1.5×10^4 /ml in EMEM medium, supplemented with 10% fetal calf serum and 1.7g/litre of sodium bicarbonate.
- 3ml aliquots of the cell suspension (per treatment) are decanted into 15ml centrifuge tubes. The cells are pelleted by centrifugation.
- To each tube 3ml of the appropriate treatment are added (diluted to 50 uM in the EMEM medium). Also included are appropriate vehicle controls, a positive control
- 25 (87MEM1 diluted to 100 ug/ml) and an isotype control (IgG2a diluted to 100 ug/ml). Incubate at 37°C for 30 mins.
- 0.5ml aliquots of the cells are seeded onto sterile dentine slices in a 48-well plate and incubated at 37°C for 2 hours. Each treatment is screened in quadruplicate.
- The slices are washed in six changes of warm PBS (10 ml / well in a 6-well plate) and
- 30 then placed into fresh treatment or control. Incubate at 37°C for 48 hours.
- tartrate resistant acid phosphatase (trap) procedure** (selective stain for cells of the osteoclast lineage).
- The slices are washed in phosphate buffered saline and fixed in 2% gluteraldehyde (in 0.2M sodium cacodylate) for 5 mins.
- 35 • They are washed in water and incubated in TRAP buffer for 5 mins at 37°C.
- Following a wash in cold water they are incubated in cold acetate buffer / fast red garnet for 5 mins at 4°C.

- Excess buffer is aspirated, and the slices are air dried following a wash in water.
- The TRAP positive osteoclasts are enumerated by bright-field microscopy and are then removed from the surface of the dentine by sonication.
- Pit volumes are determined using the Nikon/Lasertec ILM21W confocal microscope.

5

INHIBITION OF RGD-MEDIATED GPIIb-IIIa BINDING

Purification of GPIIb-IIIa

Ten units of outdated, washed human platelets (obtained from Red Cross) were lysed by gentle stirring in 3% octylglucoside, 20 mM Tris-HCl, pH 7.4, 140 mM NaCl, 2 mM CaCl_2 at 4°C for 2 h. The lysate was centrifuged at 100,000g for 1 h. The supernatant obtained was applied to a 5 mL lentil lectin sepharose 4B column (E.Y. Labs) preequilibrated with 20 mM Tris-HCl, pH 7.4, 100 mM NaCl, 2 mM CaCl_2 , 1% octylglucoside (buffer A). After 2 h incubation, the column was washed with 50 mL cold buffer A. The lectin-retained GPIIb-IIIa was eluted with buffer A containing 10% dextrose. All procedures were performed at 4°C. The GPIIb-IIIa obtained was >95% pure as shown by SDS polyacrylamide gel electrophoresis.

15

Incorporation of GPIIb-IIIa in Liposomes.

A mixture of phosphatidylserine (70%) and phosphatidylcholine (30%) (Avanti Polar Lipids) were dried to the walls of a glass tube under a stream of nitrogen. Purified GPIIb-IIIa was diluted to a final concentration of 0.5 mg/mL and mixed with the phospholipids in a protein:phospholipid ratio of 1:3 (w:w). The mixture was resuspended and sonicated in a bath sonicator for 5 min. The mixture was then dialyzed overnight using 12,000-14,000 molecular weight cutoff dialysis tubing against a 1000-fold excess of 50 mM Tris-HCl, pH 7.4, 100 mM NaCl, 2 mM CaCl_2 (with 2 changes). The GPIIb-IIIa-containing liposomes were centrifuged at 12,000g for 15 min and resuspended in the dialysis buffer at a final protein concentration of approximately 1 mg/mL. The liposomes were stored at -70°C until needed.

25

Competitive Binding to GPIIb-IIIa

The binding to the fibrinogen receptor (GPIIb-IIIa) was assayed by an indirect competitive binding method using [^3H]-SK&F-107260 as an RGD-type ligand. The binding assay was performed in a 96-well filtration plate assembly (Millipore Corporation, Bedford, MA) using 0.22 μm hydrophilic durapore membranes. The wells were precoated with 0.2 mL of 10 $\mu\text{g/mL}$ polylysine (Sigma Chemical Co., St. Louis, MO.) at room temperature for 1 h to block nonspecific binding. Various concentrations of unlabeled benzodiazepines were added to the wells in quadruplicate. [^3H]-SK&F-

35

107260 was applied to each well at a final concentration of 4.5 nM, followed by the addition of 1 µg of the purified platelet GPIIb-IIIa-containing liposomes. The mixtures were incubated for 1 h at room temperature. The GPIIb-IIIa-bound [3H]-SK&F-107260 was separated from the unbound by filtration using a Millipore filtration manifold, followed by washing with ice-cold buffer (2 times, each 0.2 mL). Bound radioactivity remaining on the filters was counted in 1.5 mL Ready Solve (Beckman Instruments, Fullerton, CA) in a Beckman Liquid Scintillation Counter (Model LS6800), with 40% efficiency. Nonspecific binding was determined in the presence of 2 µM unlabeled SK&F-107260 and was consistently less than 0.14% of the total radioactivity added to the samples. All data points are the mean of quadruplicate determinations.

Competition binding data were analyzed by a nonlinear least-squares curve fitting procedure. This method provides the IC₅₀ of the antagonists (concentration of the antagonist which inhibits specific binding of [3H]-SK&F-107260 by 50% at equilibrium). The IC₅₀ is related to the equilibrium dissociation constant (K_i) of the antagonist based on the Cheng and Prusoff equation: $K_i = IC_{50} / (1 + L / K_d)$, where L is the concentration of [3H]-SK&F-107260 used in the competitive binding assay (4.5 nM), and K_d is the dissociation constant of [3H]-SK&F-107260 which is 4.5 nM as determined by Scatchard analysis.

Preferred compounds of this invention have an affinity for the vitronectin receptor relative to the fibrinogen receptor of greater than 4:1. More preferred compounds have a ratio of activity of greater than 10:1.

Vascular smooth muscle cell migration assay

The compounds of the instant invention were tested for their ability to inhibit the migration and proliferation of smooth muscle tissue in an artery or vein in order to assess their ability to prevent restenosis of an artery, such as that which typically occurs following angioplasty.

Rat or human aortic smooth muscle cells were used. The cell migration was monitored in a Transwell cell culture chamber by using a polycarbonate membrane with pores of 8 µm (Costar). The lower surface of the filter was coated with vitronectin. Cells were suspended in DMEM supplemented with 0.2% bovine serum albumin at a concentration of $2.5 - 5.0 \times 10^6$ cells/mL, and were pretreated with test compound at various concentrations for 20 min at 20°C. The solvent alone was used as control. 0.2 mL of the cell suspension was placed in the upper compartment of the chamber. The lower compartment contained 0.6 mL of DMEM supplemented with 0.2% bovine serum albumin. Incubation was carried out at 37°C in an atmosphere of 95% air/5% CO₂ for 24 hr. After incubation, the non-migrated cells on the upper surface of the filter were

removed by gentle scraping. The filter was then fixed in methanol and stained with 10% Giemsa stain. Migration was measured either by a) counting the number of cells that had migrated to the lower surface of the filter or by b) extracting the stained cells with 10% acetic acid followed by determining the absorbance at 600 nM.

5

Examples

Nuclear magnetic resonance spectra were recorded at either 250 or 400 MHz using, respectively, a Bruker AM 250 or Bruker AC 400 spectrometer. CDCl_3 is deuteriochloroform, $\text{DMSO}-d_6$ is hexadeuteriodimethylsulfoxide, and CD_3OD is tetradeuteriomethanol. Chemical shifts are reported in parts per million (δ) downfield from the internal standard tetramethylsilane. Abbreviations for NMR data are as follows: s=singlet, d=doublet, t=triplet, q=quartet, m=multiplet, dd=doublet of doublets, dt=doublet of triplets, app=apparent, br=broad. J indicates the NMR coupling constant measured in Hertz. Infrared (IR) spectra were recorded on a Perkin-Elmer 683 infrared spectrometer in transmission mode. IR band positions are reported in inverse wavenumbers (cm^{-1}). Mass spectra were taken on either VG 70 FE, PE Syx API III, or VG ZAB HF instruments, using fast atom bombardment (FAB) or electrospray (ES) ionization techniques. Elemental analyses were obtained using a Perkin-Elmer 240C elemental analyzer. Melting points were taken on a Thomas-Hoover melting point apparatus and are uncorrected. All temperatures are reported in degrees Celsius.

Analtech Silica Gel GF and E. Merck Silica Gel 60 F-254 thin layer plates were used for thin layer chromatography. Both flash and gravity chromatography were carried out on E. Merck Kieselgel 60 (230-400 mesh) silica gel. Analytical and preparative HPLC were carried out on Rainin or Beckman chromatographs. ODS refers to an octadecylsilyl derivatized silica gel chromatographic support. 5 μ Apex-ODS indicates an octadecylsilyl derivatized silica gel chromatographic support having a nominal particle size of 5 μ , made by Jones Chromatography, Littleton, Colorado. YMC ODS-AQ® is an ODS chromatographic support and is a registered trademark of YMC Co. Ltd., Kyoto, Japan. PRP-1® is a polymeric (styrene-divinylbenzene) chromatographic support, and is a registered trademark of Hamilton Co., Reno, Nevada. Celite® is a filter aid composed of acid-washed diatomaceous silica, and is a registered trademark of Manville Corp., Denver, Colorado.

Methyl (\pm)-7-carboxy-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetate and methyl (\pm)-7-carboxy-2,3,4,5-tetrahydro-3-oxo-4-phenylethyl-1H-1,4-benzodiazepine-2-acetate was prepared by the method of Bondinell et al. WO 93/00095. Tert-butyl 3-(bromomethyl)-4-fluorobenzoate and methyl (S)-7-carboxy-2,3,4,5-

tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetate was prepared by the method of Bondinell et al. WO 95/18619.

PREPARATION OF INTERMEDIATE COMPOUNDS

5

Preparation A

Preparation of Benzyl 3-[3,4-dihydro-8-carboxy-1-methyl-2,5-dioxo-1H-1,4-benzodiazepine]-4-propanoate

10

a) 4-Iodo-2-amino benzoic acid

Oxidation of 4-iodo-2-nitrotoluene according to Sasson, et. al., J. Org. Chem. 1986, 51, 2880-83, to give 4-iodo-2-nitro benzoic acid followed by reduction of the nitro group using iron and acetic acid gives the title compound.

15

b) 7-Iodoisatoic anhydride

To a mechanically stirred ice cold solution of the compound of Preparation A(a) (26.3 g, 0.1 mol), Na₂CO₃ (10.6 g, 0.1 mol) and H₂O (250 mL), is slowly added, via an addition funnel, a solution of 1.93M COCl₂ in toluene (80 mL). After 2 h, the precipitated product is isolated by filtration, and the solid is washed successively with H₂O (200 mL), a 1:1 mixture of EtOH:Et₂O (300 mL), and Et₂O (200 mL), and dried under vacuum to yield the title compound.

20

c) Benzyl N-(2-amino-4-iodobenzoyl)-β-alanine

A magnetically stirred solution of the compound of Preparation A(b) (5.0 g, 0.0173 mol), β-alanine benzyl ester tosylate (5.85 g, 0.0173 mol), and DMAP (0.5 g, 0.0041 mol) in pyridine (35 mL) is heated for 2 h at 80° C. The reaction mixture is allowed to cool to RT and concentrated. The resulting residue is dissolved in EtOAc (100 mL), and washed successively with 10% cupric sulfate (2 x 50 mL), saturated NaHCO₃ (1 x 50 mL) and brine (1 x 50 mL), dried (Na₂SO₄), filtered, and concentrated to afford the title compound after chromatography (silica gel, 1:1 EtOAc/hexanes).

30

d) Benzyl N-(2-methylamino-4-iodobenzoyl)- β -alanine

A magnetically stirred solution of the compound of Preparation A(c) (2.0 mmol), 2,6-lutidine (0.35 mL, 3.0 mmol) and CH₃I (0.19 mL, 3.0 mmol) in DMF (15 mL) is heated at 50°C for 15 h. The reaction mixture is allowed to cool to RT and concentrated.

- 5 The resulting residue is dissolved in EtOAc (75 mL), and washed successively with 10% citric acid (1 x 50 mL), saturated NaHCO₃ (1 x 50 mL) and brine (1 x 50 mL), dried (Na₂SO₄), filtered, and concentrated to afford the title compound after chromatography (silica gel, gradient, 35-65% EtOAc/hexanes).

10 e) Benzyl 3-[3,4-dihydro-8-iodo-1-methyl-2,5-dioxo-1H-1,4-benzodiazepine]-4-propanoate

To a cold (-30°C) magnetically stirred solution of the compound of Preparation A(d) (0.305 g, 0.69 mmol), Et₃N (0.144 g, 1.04 mmol) in CH₂Cl₂ (3 mL) is added slowly a solution of α -bromoacetyl bromide (0.09 mL, 1.04 mmol) in CH₂Cl₂ (2 mL)

- 15 under argon atmosphere. The reaction mixture is allowed to warm to RT and stir for 2 h. The mixture is diluted with CH₂Cl₂ (40 mL) and washed successively with 10% citric acid (1 x 50 mL), saturated NaHCO₃ (1 x 50 mL), dried (Na₂SO₄), filtered, and concentrated. The resulting residue is dissolved in DMF (3 mL) and added via an addition funnel to a slurry of NaH (25 mg, 1.04 mmol) in DMF (2 mL) which is cooled to
20 0°C. After 2 h of stirring, the mixture is poured into an ice cold solution of 10% citric acid (50 mL) and extracted with EtOAc (3 x 40 mL). The combined extracts are washed with saturated NaHCO₃ (1 x 50 mL), dried (Na₂SO₄), filtered, and concentrated to afford the title compound after chromatography (silica gel, gradient, 40-70% EtOAc/hexanes).

25 f) Benzyl 3-[3,4-dihydro-8-carboxy-1-methyl-2,5-dioxo-1H-1,4-benzodiazepine]-4-propanoate

A solution of the compound of Preparation A(e) (3.2 mmol), Pd(OAc)₂ (0.16 mmol), and 1,1'-bis(diphenylphosphine)ferrocene (0.64 mmol,) in DMSO (20 mL) is heated to 65°C under a carbon monoxide balloon for 18 h. The reaction mixture is
30 diluted with water, acidified with 1 N HCl and extracted with CH₂Cl₂. The combined

organic extracts are washed with water, dried (Na₂SO₄), filtered, and concentrated to afford the title compound after chromatography (silica gel).

Preparation B

5

Ethyl 3-[4H-imidazo[1,2-a][1,4]benzodiazepine-5(6H)-1-methyl-6-oxo-9-carboxyl]-5-propanoic acid

a) Ethyl N-(2-amino-4-iodobenzoyl)-β-alanine

10 A magnetically stirred solution of the compound of Preparation A(b) (0.0173 mol), β-alanine ethyl ester hydrochloride (0.0173 mol), and DMAP (0.5 g, 0.0041 mol) in pyridine (35 mL) is heated for 2 h at 80°C. The reaction mixture is allowed to cool to RT and concentrated. The resulting residue is dissolved in EtOAc (100 mL), and washed successively with 10% cupric sulfate (2 x 50 mL), saturated NaHCO₃ (1 x 50 mL) and
15 brine (1 x 50 mL), dried (Na₂SO₄), filtered, and concentrated to afford the title compound after chromatography (silica gel, 1:1 EtOAc/hexanes).

b) Ethyl 3-[3,4-dihydro-8-iodo-2,5-dioxo-1H-1,4-benzodiazepine]-4-propanoate

To a cold (-30°C) magnetically stirred solution of the compound of Preparation
20 B(a) (0.69 mmol), and Et₃N (0.144 g, 1.04 mmol) in CH₂Cl₂ (3 mL) is added slowly a solution of α-bromoacetyl bromide (0.09 mL, 1.04 mmol) in CH₂Cl₂ (2 mL) under argon atmosphere. The reaction mixture is allowed to warm to RT and stir for 2 h. The mixture is diluted with CH₂Cl₂ (40 mL) and wash successively with 10% citric acid (1 x 50 mL), saturated NaHCO₃ (1 x 50 mL), dried (Na₂SO₄), filtered, and concentrated. The
25 resulting residue is dissolved in DMF (3 mL) and added via an addition funnel to a slurry of NaH (25 mg, 1.04 mmol) in DMF (2 mL) which is cooled to 0°C. After 2 h of stirring, the mixture is poured into an ice cold solution of 10% citric acid (50 mL) and extracted with EtOAc (3 x 40 mL). The combined extracts are washed with saturated NaHCO₃ (1 x 50 mL), dried (Na₂SO₄), filtered, and concentrated to afford the title compound after
30 chromatography (silica gel).

c) Ethyl-3-[3,4-dihydro-8-iodo-2-thioxo-5-oxo-1H-1,4-benzodiazepine]-4-propanoate

To a solution of the compound of Preparation B(b) (1.0 g, 2.49 mmol) in THF (10 mL) at RT and under an atmosphere of nitrogen is added Lawesson's reagent (1.0 g) and the reaction is heated at 50°C for 2 h. The reaction mixture is allowed to cool to RT and is concentrated. Purifying the resulting residue by chromatography (silica gel, gradient, 40-60% EtOAc/hexane) gives the title compound.

d) Ethyl 3-[4H-imidazo[1,2-a][1,4]benzodiazepine-5(6H)-1-methyl-6-oxo-9-iodo]-5-propanoate

To a vigorously stirred biphasic solution of the compound of Preparation B(c) (0.95 g, 2.27 mmol), CH₃I (0.2 g) and a catalytic amount of tetrabutylammonium hydrogen sulfate in CH₂Cl₂ (10 mL) and H₂O (10 mL), is added 2 N NaOH (1.2 mL) at RT. After 2 h, the layers are separated and the aqueous layer is washed with CH₂Cl₂ (2 x 25 mL). The combined organic extracts are dried (Na₂SO₄), filtered, and concentrated. The resulting residue is dissolved in toluene (10 mL) and allowed to react with propargyl amine (0.64 mL) and pyridine hydrochloride (0.23 g). The reaction is heated to reflux for 6 h, allowed to cool to RT, and concentrated to give the title compound after chromatography (silica gel, EtOAc).

e) Ethyl 3-[4H-imidazo[1,2-a][1,4]benzodiazepine-5(6H)-1-methyl-6-oxo-9-carboxy]-5-propanoic acid

A solution of the compound of Preparation B(d) (3.2 mmol), Pd(OAc)₂ (0.16 mmol), and 1,1'-bis(diphenylphosphine)ferrocene (0.64 mmol) in DMSO (20 mL) is heated at 65°C under a carbon monoxide balloon for 18 h. The reaction mixture is diluted with H₂O, acidified with 1 N HCl and extracted with CH₂Cl₂ (3 x). The combined organic extracts are washed with H₂O, dried (Na₂SO₄), filtered, and concentrated to afford the title compound after chromatography (silica gel).

Preparation C

Preparation of Ethyl 4-(1-piperazinyl)-1-piperidineacetate

- 5 a) Ethyl 4-[4-(tert-butoxycarbonyl)-1-piperazinyl]-1-piperidineacetate

The titled compound is prepared from tert-butyl 1-piperazinecarboxylate (Aldrich) and ethyl 4-oxo-1-piperidineacetate (Porter, et. al., EP 0 542 363 A2) by reductive amination with NaBH₃CN according to the method of Porter, et. al., EP 0 542 363 A2.

- 10 b) Ethyl 4-(1-piperazinyl)-1-piperidineacetate

A solution of Preparation C(a) and 4M HCl in dioxane/CH₂Cl₂ is stirred at RT for 18 h. The reaction mixture is concentrated to give the title compound as the hydrochloride salt.

15

Preparation D

Preparation of 6-Methyl-2-(phthalimido)pyridine

A mixture of 6-methyl-2-aminopyridine and neat phthalic anhydride is heated for 5 h, diluted with aqueous NaHCO₃, and extracted with EtOAc. The combined extracts

- 20 are dried and evaporated to give the title compound.

Preparation E

Preparation of 3-(6-Amino-2-pyridinyl)propanol

25

- a) 1-[6-(3-Hydroxypropyl)-2-pyridinyl]-2,5-dimethylpyrrole

Following the procedure of Warter, et. al., Org. Synth. **1943**, 23, 83, for the preparation of 2-(3-hydroxypropyl)pyridine, except substituting 1-(6-methylpyridin-2-yl)-2,5-dimethylpyrrole for the 2-methylpyridine, J. Chem. Soc., Perkin Trans. 1, **1984**,

- 30 2801-2807, gives the title compound.

b) 3-(6-Amino-2-pyridinyl)propanol

Following the procedure of Bruckelman, et. al., J. Chem. Soc. Perkin Trans. 1, 1984, 2801-2807, for the preparation of 2-amino-6-ethylpyridine, except substituting the compound of Preparation E(a) for 1-(6-ethylpyridin-2-yl)-2,5-dimethylpyrrole, gives the title compound.

Preparation FPreparation of 4-[6-(Toluenesulfonylamino)-2-pyridinyl]-1-propanol oxime

a) 4-[6-(2,5-Dimethylpyrrol-1-yl)-2-pyridinyl]-1-butene

Following the procedure of Meakins, J. Chem. Soc. Perkin Trans. I, 1984, 2801, for the alkylation of 6-(2,5-dimethylpyrrol-1-yl)-2-picoline, except using allyl bromide as the alkylating agent, the title compound is prepared.

b) 4-(6-Amino-2-pyridinyl)-1-butene

Following Procedure B, described Meakins, J. Chem. Soc. Perkin Trans. I, 1984, 2801, the compound of Preparation F(a) is deprotected to give the title compound.

c) 4-[6-(Toluenesulfonylamino)-2-pyridinyl]-1-butene

Sodium hydride (55 mmol) is added carefully to a solution of the compound of Preparation F(b) (50 mmol) and 4-toluenesulfonyl chloride (55 mmol) in dry THF (200 mL). The reaction is stirred at RT until complete, then is quenched with saturated NH_4Cl (200 mL), and the mixture is extracted with EtOAc. The combined organic extracts are dried (MgSO_4) and concentrated, and the residue is purified by chromatography (silica gel) to give the title compound.

d) 4-[6-(Toluenesulfonylamino)-2-pyridinyl]-1-propanal

Ozone is bubbled into a solution the compound of Preparation F(c) (40 mmol) in CH_2Cl_2 (160 mL) and CH_3OH (40 mL) at -78°C until the blue color persists, then the excess ozone is removed by bubbling argon through the solution. Dry dimethylsulfide

(excess) is added, and the reaction is warmed to RT. The reaction is stirred at RT until complete, then is concentrated, and the residue is chromatographed (silica gel) to afford the title compound.

5 e) 4-[6-(Toluenesulfonylamino)-2-pyridinyl]-1-propanal oxime

Hydroxylamine hydrochloride (33 mmol) is added to a solution of the compound of Preparation F(d) (30 mmol) and anhydrous NaOAc (66 mmol) in CH₃OH (150 mL) at 0°C. The reaction is stirred at 0°C until complete, then is concentrated, and the residue is partitioned between H₂O and EtOAc. The layers are separated, and the aqueous layer is
10 extracted with EtOAc. The combined organic layers are washed sequentially with 5% NaHCO₃ and saturated brine, dried (MgSO₄), and concentrated to afford the title compound.

Preparation G

15

Preparation of (6-Amino-2-pyridinyl)acetic acid

Ethyl (6-amino-2-pyridinyl)acetate, Awaya, et. al., Chem. Pharm. Bull., 1974, 22, 1414, (1 mmol) is treated with 1N NaOH (1.5 mmol) in CH₃OH (20 mL). The mixture is concentrated, extracted with CH₂Cl₂, and the aqueous phase is adjusted to pH 5 to give
20 the title compound.

Preparation H

Preparation of 6-(2-Aminoethyl)-2-pyridinamine dihydrochloride

25 Following the procedures of Preparation 13 in Bondinell, et al., WO 94/14476, for the preparation of 2-aminopyridine-4-ethanamine dihydrochloride, except substituting 2-(acetylamino)pyridine-6-carboxylic acid for the 2-(acetylamino)pyridine-4-carboxylic acid, the title compound is prepared.

Preparation I

Preparation of 4-[(6-Amino-2-pyridinyl)methyl]phenol

4-[(6-Amino-2-pyridinyl)methyl]anisole, Ife et. al., WO 9426715, is heated with
5 concentrated hydrobromic acid to afford the title compound.

Preparation J

Preparation of Benzyl 4-[2-(methylamino)acetyl]phenoxyacetate hydrochloride

10

a) 4-[N-Boc-2-(methylamino)acetyl]phenol

A solution of di-tert-butyl dicarbonate (5.96 g, 27.3 mmol) in 1,4-dioxane (25 mL) was added dropwise at 0°C to a mixture of 4-[2-(methylamino)acetyl]phenol hydrochloride (5.0 g, 24.8 mmol), 1,4-dioxane (30 mL), H₂O (25 mL), and 1.0 N NaOH
15 (25 mL, 25 mmol). After 24 h, the reaction was warmed to RT and stirred for 1.5 h. More 1.0 N NaOH (25 mL, 25 mmol) was added, and the reaction was stirred for an additional 0.5 h at RT, and concentrated. The residue was diluted with EtOAc (80 mL), and the mixture was acidified to pH 2 using 1.0 M NaHSO₄. The resulting mixture was extracted with EtOAc, and the combined organic layers were washed with H₂O and dried
20 (Na₂SO₄). Filtration and concentration gave the title compound (6.49 g, 99%): ¹H NMR (250 MHz, CDCl₃) δ 6.70-8.05 (m, 4 H), 4.53 (s, 2H), 2.98 (s, 3H), 1.50 (s, 9H).

b) Benzyl 4-[N-Boc-2-(methylamino)acetyl]phenoxyacetate

A mixture of the compound of Preparation J(a) (5.04 g, 19.0 mmol) and K₂CO₃
25 (2.63 g, 19.0 mmol) in acetone (100 mL) was stirred at reflux under argon for 1h. The mixture was cooled to RT and benzyl bromoacetate (5.23 g, 22.8 mmol) was added. The reaction was heated at reflux for 18 h, then was cooled and filtered. The filter cake was washed with acetone, and the filtrate was concentrated. The residue was dissolved in CH₂Cl₂ (300 mL) and washed sequentially with H₂O (50 mL) and brine (50 mL).
30 Drying (Na₂SO₄), concentration, and flash chromatography (silica gel, 1:3

EtOAc/hexanes) yielded the title compound (7.28 g, 93%): ^1H NMR (250 MHz, CDCl_3) δ 6.85-7.95 (m, 9 H), 5.23 (s, 2H), 4.71 (s, 2H), 4.55 (d, 2H), 2.95 (d, 3H), 1.45 (d, 9H).

c) Benzyl 4-[2-(methylamino)acetyl]phenoxyacetate hydrochloride

- 5 A mixture of the compound of Preparation J(b) (7.26 g, 17.57 mmol) and 4 M HCl in 1,4-dioxane (150 mL) was stirred for 1 h at RT. Concentration and trituration with Et_2O afforded the title compound as a white powder (5.93 g, 97%): ^1H NMR (250 MHz, CD_3OD) δ 7.05-8.00 (m, 9 H), 5.23 (s, 2H), 4.88 (s, 2H), 4.65 (s, 2H), 2.80 (s, 3H).

10

Preparation K

Preparation of Dimethyl 4-[2-(methylamino)acetyl]-1,2-phenylenedioxydiacetate hydrochloride

15

a) 4-[N-Boc-2-(methylamino)acetyl]-1,2-dihydroxybenzene

Following the procedure of Preparation J(a), except substituting adrenalone hydrochloride (5.0 g, 23.0 mmol) for 4-[2-(methylamino)acetyl]phenol hydrochloride, the title compound (1.2 g, 19%) was prepared following flash chromatography (silica gel, 1:1

- 20 EtOAc/hexanes): MS (ES) m/e 282.2 $[\text{M}+\text{H}]^+$.

b) Dimethyl 4-[N-Boc-2-(methylamino)acetyl]-1,2-phenylenedioxydiacetate

- Following the procedure of Preparation J(b), except substituting the compound of Preparation K(a) (0.9 g, 3.2 mmol) for the compound of Preparation J(a) and methyl bromoacetate (1.23 g, 8.0 mmol) for benzyl bromoacetate, the title compound (1.11 g, 81%) was prepared: MS (ES) m/e 426.2 $[\text{M}+\text{H}]^+$.
- 25

c) Dimethyl 4-[N-Boc-2-(methylamino)acetyl]-1,2-phenylenedioxydiacetate hydrochloride

Following the procedure of Preparation J(c), except substituting the compound of Preparation K(b) (1.11 g, 2.6 mmol) for the compound of Preparation J(b), the title compound was prepared (1.1 g, quantitative): MS (ES) m/e 326.0 [M+H]⁺.

Preparation L

Preparation of (6-Phthaloyl-2-pyridinyl)methanamine

Following the procedure of Preparation T, except substituting ammonia for methylamine, gives the title compound.

Preparation M

15 Preparation of Ethyl (6-carboxy-1,2,3,4-tetrahydroisoquinolin-2-yl)acetate

a) Ethyl (6-Methoxy-1,2,3,4-tetrahydroisoquinolin-2-yl)acetate

A solution of 6-methoxy-1,2,3,4-tetrahydroisoquinoline, Sall and Grunewald, J. Med. Chem. **1987**, *30*, 2208-2216, (1.1 mmol), ethyl chloroacetate (1.17 mmol), and K₂CO₃ (1.17 mmol) in CH₃CN (10 mL) is stirred for 18 h. The mixture is partitioned between EtOAc and H₂O. The organic phase is concentrated to an oil, which is purified by chromatography (silica gel, gradient, 20-80% EtOAc/hexane) to afford the title compound.

25 b) Ethyl (6-Hydroxy-1,2,3,4-tetrahydroisoquinolin-2-yl)acetate

A solution of the compound of Preparation M(a) (0.249 g, 1.0 mmol), 1M BBr₃ in CH₂Cl₂ (1.0 mL, 1.0 mmol) is stirred at -70°C for 2 h and then stirred at RT for 12 hr. The solution is concentrated, and the solution of the resulting oil in EtOAc is washed with H₂O, 5% NaHCO₃, and H₂O, dried (MgSO₄), filtered, and concentrated to an oil to afford the title compound (0.223 g, 95%)

c) Ethyl [6-(trifluoromethylsulfonyloxy)-1,2,3,4-tetrahydroisoquinolin-2-yl]acetate

A solution of the compound of Preparation M(b) (0.235 g, 1.0 mmol), trifluorosulfonic acid anhydride (0.23 mL, 1.1 mmol) and Et₃N (0.32 mL, 1.5 mmol) in CH₂Cl₂ (5 mL) is stirred for 8 h. The solution is concentrated to an oil which is taken up in EtOAc. The organic phase is washed with 5% NaHCO₃ and H₂O. The organic phase is dried (Na₂SO₄), filtered, concentrated to afford the title compound (0.300 g, 82%)

d) Ethyl (6-carboxy-1,2,3,4-tetrahydroisoquinolin-2-yl)acetate

A solution of the compound of Preparation M(c) (0.367 g, 1.0 mmol), Pd(OAc)₂ (0.022 g, 0.1 mmol), Ph₃P (0.262 g, 1.0 mmol), diisopropylamine (0.34 mL, 2.5 mmol), and NMP (5 mL) in 10% NH₄CO₃ is stirred for 8 h under an atmosphere of CO. The solution is concentrated to an oil which is purified by chromatography (silica gel, gradient, 10-33% CH₃OH/CH₂Cl₂) to afford the title compound (0.19 g, 72%).

15

Preparation NPreparation of Ethyl (6-carboxy-1,2,3,4-tetrahydro-1-oxo-isoquinolin-2-yl)acetate

a) Ethyl (6-Methoxy-1-oxo-1,2,3,4-tetrahydroisoquinolin-2-yl)acetate

A mixture of 6-methoxy-1,2,3,4-tetrahydro-1-oxo-isoquinoline, Sall and Grunewald, J. Med. Chem., 1987, 30, 2208-2216, (0.39 mmol) and NaH (0.17 g, 0.43 mmol, 60% oil dispersion) in THF (5 mL) is heated to reflux for 1 h and then allowed to cool to RT. Ethyl chloroacetate (0.43 mmol) is added and the mixture is allowed to stir for 1 h. The mixture is quenched with H₂O (10 mL) and washed with EtOAc. The organic layers are combined, washed with H₂O (10 mL) and concentrated to an oil which is purified by (silica gel, gradient, 10-33% CH₃OH/CH₂Cl₂) to afford the title compound.

b) Ethyl (6-Hydroxy-1-oxo-1,2,3,4-tetrahydroisoquinolin-2-yl)acetate

A solution of the compound of Preparation N(a) (0.263 g, 1.0 mmol) and 1M BBr₃ in CH₂Cl₂ (1.1 mL) is stirred at -70°C for 2 h and then at RT for 4 h. The solution is concentrated to an oil which is taken up in EtOAc. The organic phase is washed with H₂O, 5% NaHCO₃, H₂O, dried (MgSO₄), filtered, and concentrated to afford the title compound (0.20 g, 80%).

c) Ethyl [6-(trifluoromethylsulfonyloxy)-1,2,3,4-tetrahydro-1-oxo-isoquinolin-2-yl]acetate

A solution of the compound of Preparation N(b) (3.4 mmol) and trifluorosulfonic acid anhydride (3.4 mmol, mL) in pyridine (5 mL) is chilled at 0°C and allowed to warm RT for 1 h. The mixture is quenched with H₂O (5 mL) and washed with EtOAc. The organic layers are combined, washed with H₂O (7 mL) and concentrated to an oil. The residue is purified chromatography (by silica gel, gradient, 14-75% EtOAc/hexane) to afford the title compound.

d) Ethyl (6-carboxy-1-oxo-1,2,3,4-tetrahydroisoquinolin-2-yl)acetate

A solution of the compound of Preparation N(c) (0.23 g, 1.0 mmol), Pd(OAc)₂ (0.026 g, 0.1 mmol), Ph₃P (0.262 g, 1.0 mmol), diisopropylamine (0.23 mL, 2.0 mmol) and NMP (7 mL) in 10% NH₄CO₃ is stirred for 8 h under an atmosphere of CO. The solution is concentrated to an oil which is purified by chromatography (silica gel, gradient, 25-75% CH₃OH/CH₂Cl₂) to afford the title compound (0.31 g, 70%).

Preparation OPreparation of Ethyl (6-carboxy-tetralin-2-yl)acetate

a) Ethyl [6-(trifluoromethylsulfonyloxy)-tetralin-2-yl]acetate

Following the procedure of Preparation M(c), except substituting ethyl (6-hydroxy-tetralin-2-yl)acetate, Fisher, et. al., EP 0635492, Scheme 6 and Example 20, parts A-D for the compound of Preparation M(b), gives the title compound.

b) Ethyl (6-carboxy-tetralin-2-yl)acetate

Following the procedure of Preparation M(d), except substituting the compound of Preparation O(a) for the compound of Preparation M(c), gives the title compound.

5

Preparation PPreparation of Ethyl (5-aminobenzofuran-2-yl)propionate and Ethyl (5-amino-2,3-dihydro-benzofuran-2-yl)propionate

10

a) 2-(Ethoxycarbonyl)methoxy-5-nitrobenzaldehyde

A solution of 5-nitrosalicylaldehyde (Aldrich) (0.167 g, 1.0 mmol), ethyl bromoacetate (0.166 g, 1.0 mmol), K_2CO_3 (0.276 g, 2.0 mmol) and NaI (0.015 g, 0.1 mmol) in THF (10 mL) is heated to 80°C for 24 h. The solution is concentrated and the residue is purified by chromatography (silica gel, gradient, 5-20% CH_3OH in CH_2Cl_2) to afford the title compound (0.20 g, 87%)

15

b) Ethyl (5-nitrobenzofuran-2-yl)carboxylate

A solution of the compound of Preparation P(a) (0.229 g, 1.0 mmol) and DBU (0.152 g, 1.0 mmol) in EtOH (10 mL) is allowed to stir at RT for 18 h. The solution is concentrated and the residue is treated with EtOH (10 mL). The solution is bubbled with HCl gas for 2 min and refluxed for 5 h. The solution is concentrated and the residue is treated with EtOAc. The organic phase is washed with H_2O , 5% citric acid, H_2O , 5% $NaHCO_3$, and H_2O . The organic phase is concentrated to afford the title compound (0.19 g, 81%).

25

c) Ethyl (5-nitrobenzofuran-2-yl)carboxaldehyde

A cold solution (-78°C) of the compound of Example 20(b) (0.235 g, 1.0 mmol) in THF (5 mL) is treated with 1M DiBAL in THF (1.0 mL, 1.0 mmol). The solution is stirred at -78°C for 30 min and at RT for 3 h. The solution is treated with CH_3CO_2H (3 mL) followed by H_2O (2 mL). The solution is concentrated and the residue is treated

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with toluene to azeotrope off the acetic acid. Drying *in vacuo* afforded the title compound (0.100 g, 52%)

d) Ethyl (5-nitrobenzofuran-2-yl)propenoate

5 A solution of triethyl phosphonoacetate (0.224 g, 1.0 mmol) in THF (5 mL) is treated with NaH (60% suspension in mineral oil, 0.04 g, 1.0 mmol) at 0° C for 1 h. To the solution is added the compound of Preparation P(c) (0.235 g, 1.0 mmol). The solution is stirred at RT for 18 h, concentrated, and the residue is purified by chromatography (silica gel, gradient, 5-20% EtOAc/hexane) (EtOAc/Hexane 0.5:9 to 4:1) to afford the
10 title compound (0.2g, 77%).

e) Ethyl (5-aminobenzofuran-2-yl)propionate and Ethyl (5-amino-2,3-dihydrobenzofuran-2-yl)propionate

 A solution of the compound of Preparation P(d) (0.261 g, 1.0 mmol) in EtOH (5
15 mL) containing 10% Pd/C (0.026 g) is hydrogenated at 45 psi for 1 h. The solution is filtered through Celite and the filtrate is concentrated and chromatographed (silica gel, gradient, 25-75% EtOAc/hexane) affords the title compounds.

Preparation Q

20

Preparation of Ethyl (5-carboxy-benzofuran-2-yl)propionate

a) Ethyl [5-(tert-butyldimethylsilyloxy)benzofuran-2-yl]carboxylate

 A solution of ethyl [5-(hydroxy)benzofuran-2-yl]carboxylate, Denny, et. al., EP
25 0655439, (0.206 g, 1.0 mmol), tert-(butyl)dimethylsilyl chloride (0.23 mL, 1.0 mmol) and imidazole (0.34 g, 1.0 mmol) in THF is allowed to stir for 4 h. The solution is concentrated and the residue is treated with EtOAc. The organic phase is washed with H₂O, dried (Na₂SO₄), and concentrated to afford the title compound (0.35 g, 90%).

30 b) Ethyl [5-[tert-(butyl)dimethylsilyloxy]benzofuran-2-yl]propenoate

Following the procedure of Preparation P(c) and (d), except substituting the compound of Preparation Q(a) for the compound of Preparation P(b), gives the title compound.

- 5 c) Ethyl [5-(hydroxy)benzofuran-2-yl]propionate and Ethyl [5-hydroxy-2,3-dihydro-benzofuran-2-yl]propionate

A mixture of the compound of Preparation Q(b) (0.234g, 1.2 mmol) and 10% Pd/C (0.023 g, 10% wt) in EtOH(5 mL) is hydrogenated at 50 psi for 1 h. The mixture is filtered through Celite and concentrated. A solution of the residue (0.34 g, 1.0 mmol) and 10 Et₄NF (0.149 g, 1.0 mmol) in THF (10 mL) is allowed to stir at RT for 18 h. The solution is concentrated and purified by chromatography (silica gel) to give the title compounds (0.25 g, 57%).

- d) Ethyl [5-(trifluoromethylsulfonyloxy)benzofuran-2-yl]propionate

15 Following the procedure of Preparation M(c), except substituting ethyl [5-(hydroxy)benzofuran-2-yl]propionate of Preparation Q(c) for the compound of Preparation M(b), gives the title compound.

- e) Ethyl (5-carboxy-benzofuran-2-yl)propionate

20 Following the procedure of Preparation M(d), except substituting the compound of Preparation Q(d) for the compound of Preparation M(c), gives the title compound.

Preparation R

- 25 Preparation of Ethyl (5-carboxy-2,3-dihydro-benzofuran-2-yl)propionate

- a) Ethyl [5-(trifluoromethylsulfonyloxy)-2,3-dihydro-benzofuran-2-yl]propionate

Following the procedure of Preparation Q(d), except substituting ethyl [5-hydroxy-2,3-dihydro-benzofuran-2-yl]propionate from Preparation Q(c) for ethyl [5-
30 (hydroxy)benzofuran-2-yl]propionate from Preparation Q(c), gives the title compound.

b) Ethyl (5-carboxy-2,3-dihydro-benzofuran-2-yl)propionate

Following the procedure of Preparation Q(e), except substituting the compound of Preparation R(a) for the compound of Preparation Q(d), gives the title compound.

5

Preparation SPreparation of Ethyl (±)-3-[(glycyl)amino]-4-pentynoate trifluoroacetate

10 a) Ethyl (±)-3-[[[N-tert-butoxycarbonyl]glycyl]amino]-4-pentanoate

DIEA (0.92 mL, 5.32 mmol) was added to a stirred solution of ethyl (±)-3-amino-4-pentynoate (0.3 g, 2.13 mmol), Boc-Gly (0.56 g, 3.19 mmol), HOBt · H₂O (0.43 g, 3.19 mmol), and EDC (0.61 g, 3.19 mmol) in anhydrous CH₃CN (15 mL) at RT. After 34 h, the reaction mixture was concentrated, diluted with CH₂Cl₂ (70 mL), and washed sequentially with 5% NaHCO₃ (2x15 mL) and brine (15 mL). Drying (MgSO₄), concentration, and chromatography (silica gel, 1:1 EtOAc/hexane) gave the title compound (0.5 g, 79%) as a colorless oil: MS (ES) m/e 299.2 (M+H)⁺.

20 b) Ethyl (±)-3-[(glycyl)amino]-4-pentynoate trifluoroacetate

A solution of TFA (5 mL) and CH₂Cl₂ (15 mL) at RT was added all at once to the compound of Preparation S(a) (0.5 g, 1.68 mmol). After 30 min, the solution was concentrated, and the residue was reconstituted from toluene (to remove residual TFA) to afford the title compound (0.55g, 106%) as a light yellow syrup: MS (ES) m/e 199.2 (M+H)⁺.

25

Preparation TPreparation of 6-(Methylamino)methyl-2-pyridinamine

6-Bromomethyl-2-(phthalimido)pyridine (prepared according to the method of US 4,490,533)(1.1 g, 3 mmol) was added to a solution of ethanol (100 mL) saturated with methylamine at 0°C. The resulting solution was stirred at 0°C for 2 h, concentrated to a volume of 20 mL, and treated with hydrazine hydrate (1 mL, 20 mmol). The resulting solution was heated to reflux for 2 h, concentrated, and the residue was chromatographed

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(silica gel; step gradient, 5%-15% CH₃OH/CH₂Cl₂) to give the title compound as a yellow oil (0.15 g, 32%): ¹H NMR (400 MHz, DMSO-d₆) δ 7.34 (t, J=7.8 Hz, 1H), 6.55 (d, J=7.1 Hz, 1H), 6.47 (d, J=8.3 Hz, 1H), 3.93 (s, 3H), 2.61 (s, 3H).

5

Preparation UPreparation of 6-Aminomethyl-2-pyridinamine

a) 2-(Phthalimido)methyl-6-(phthalimido)pyridine

10 A mixture of 6-bromomethyl-2-(phthalimido)pyridine (US 4,490,533)(0.4 g, 1.2 mmol), potassium phthalimide (0.30 g, 1.6 mmol), and DMF (4 mL) was stirred at RT for 18 h, concentrated, and the residue was partitioned between EtOAc and H₂O. The organic phase was washed with brine, dried (MgSO₄), and concentrated. The residue was recrystallized (CHCl₃) to give the title compound as a white solid (0.4 g, 83%): MS (ES) m/e 383.9 [M+H]⁺.

15

b) 6-Aminomethyl-2-pyridinamine

A solution of the compound of Preparation U(a)(0.4 g) and hydrazine hydrate (2 mL) in ethanol (10 mL) was heated to reflux for 2 h, filtered, and the filtrate was concentrated. The residue was triturated with CHCl₃, and the organic extracts were combined and concentrated to give the title compound as an amber oil (0.09 g, 70%): MS (ES) m/e 123.7 [M+H]⁺.

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Preparation V

25

Preparation of N-Ethyl-6-(aminomethyl)-2-pyridinamine

6-(Acetylamino)picolinamide (Farmaco Ed. Sci., 1959, 14, 594) (0.3 g, 1.67 mmol) suspended in dry THF (5 mL) was added dropwise to a solution of 1M LAH in THF (16.7 mL) cooled to 0°C. The suspension was allowed to warm to RT and was heated to reflux for 4 h. The mixture was cooled, carefully treated with H₂O and 10% NaOH, and filtered. The filtrate was dried (MgSO₄), concentrated, and the residue was azeotroped several times with toluene. The resulting mixture was concentrated to give the title compound (0.25 g, 99%): MS (ES) m/e 152 [M+H]⁺.

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Preparation WPreparation of 4-(Methylaminomethyl)-2-pyrimidinamine

A suspension of NaOAc (0.39 g, 4.8 mmol) in a solution of methylamine hydrochloride (0.39 g, 5.8 mmol) in EtOH (30 mL) was stirred 10 min at RT and 4-formyl-2-pyrimidinamine (obtained by the method of WO 9502591) (0.30 g, 2.4 mmol) was added in one portion. The mixture was stirred 30 min and NaBH₃CN (0.09 g, 1.44 mmol) was added. The resulting suspension was stirred 16 h, filtered, and concentrated. The residue was partitioned between CH₂Cl₂ (150 mL) and 5% Na₂CO₃ (20 mL). The aqueous layer was extracted with CH₂Cl₂ (4 x 25 mL), and the combined organic layers were dried (K₂CO₃) and concentrated to give the crude title compound (0.40 g): MS (ES) m/e 139 [M+H]⁺.

EXAMPLES

Example 1Preparation of (S)-7-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid

a) Methyl (S)-7-[[[(6-amino-2-pyridinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetate

EDC (0.25 g, 1.3 mmol) was added to a solution of methyl (S)-7-carboxy-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetate (0.32 g, 1.1 mmol), the compound of Preparation T (0.15 g, 6.02 mmol), HOBT·H₂O (170 mg, 1.3 mmol), and DIEA (0.9 mL, 4.4 mmol) in anhydrous CH₃CN (5 mL) at RT. After 21 h, the reaction was concentrated and the residue was partitioned between EtOAc and H₂O. The organic layer was washed with brine, dried (MgSO₄), and concentrated. The residue was purified by chromatography (silica gel, step gradient, 2%-7% CH₃OH/CH₂Cl₂) to give the title compound (0.22 g, 48%): MS (ES) m/e 412.4 [M+H]⁺.

b) (S)-7-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid

A solution of the compound of Example 1(a) (0.22 g, 54 mmol), LiOH·H₂O (0.033 g, 0.79 mmol), THF (5 mL), and water (2 mL) was stirred at RT overnight. The mixture was concentrated and the residue was dissolved in water. The resulting solution was brought to pH 5 with 3N HCl and allowed to stand. The crystals that formed were

collected by filtration and dried to give the title compound as a pale yellow solid (0.125 g, 59%): MS (ES) m/e 398.4 [M+H]⁺. Anal. Calcd for C₂₀H₂₃N₅O₄ · 3/8 H₂O: C, 59.43; H, 5.92; N, 17.33. Found: C, 59.42; H, 5.73; N, 17.18.

5

Example 2Preparation of (S)-7-[[[(6-Amino-2-pyridinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid

- 10 a) Methyl (S)-7-[[[(6-amino-2-pyridinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetate

Following the procedure of Example 1(a), except substituting the compound of Preparation U(b) for the compound of Preparation T, gave the title compound as a white foam: MS (ES) m/e 398.0 [M+H]⁺.

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- b) (S)-7-[[[(6-Amino-2-pyridyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid

Following the procedure of Example 1(b), except substituting the compound of Example 2(a) for the compound of Example 1(a), gave the title compound as a white solid: MS (ES) m/e 384.2 [M+H]⁺. Anal. Calcd. for C₁₉H₂₁N₅O₄ · 1.25 H₂O: C, 56.22; H, 5.83; N, 17.25. Found: C, 56.01; H, 5.99; N, 16.92.

20

Example 3

- 25 Preparation of (S)-7-[[[(6-Ethylamino-2-pyridinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid

- a) Methyl (S)-7-[[[(6-ethylamino-2-pyridinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetate

30

A mixture of the compound of Preparation V (0.25 g, 1.65 mmol), methyl (S)-7-carboxy-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetate (0.58 g, 2 mmol), EDC (0.38 g, 2 mmol), and HOBT·H₂O (0.26 g, 2 mmol) in DMF (20 mL) was stirred at RT overnight. The mixture was concentrated, and the residue was treated with 5% Na₂CO₃ and extracted with CH₂Cl₂ (3 x 30 mL). The combined organic extracts were washed with H₂O, dried (MgSO₄), concentrated. The residue was chromatographed (silica gel, 5% CH₃OH/CH₂Cl₂) to give the title compound (0.16 g, 23%): MS (ES) m/e 426 [M+H]⁺.

35

b) (S)-7-[[[(6-Ethylamino-2-pyriinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid

The compound of Example 3(a) (0.16 g, 0.4 mmol) was dissolved in CH₃OH (10 mL) and THF (1 mL), and treated with 1N NaOH (0.5 mL). The mixture was stirred overnight, concentrated, and the residue was dissolved in H₂O and extracted with CH₂Cl₂. The pH of the aqueous phase was adjusted to 5.5-6 with dilute HCl, and the solid which formed was filtered, washed with H₂O and Et₂O, and dried to give the title compound (0.11 g, 73%): MS (ES) m/e 412 [M+H]⁺. Anal. Calcd for C₂₁H₂₅N₅O₄ · 0.625 H₂O: C, 59.91; H, 6.08; N, 16.21. Found: C, 59.67; H, 6.26; N, 16.51.

Example 4

Preparation of (±)-7-[[[(2-Amino-4-pyrimidinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid

a) Methyl (±)-7-[[[(2-amino-4-pyrimidinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1,4-benzodiazepine-2-acetate

A solution methyl (±)-7-(chlorocarbonyl)-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetate hydrochloride (0.43 g, 1.25 mmol) in CH₂Cl₂ (45 mL) was added dropwise to a solution of the compound of Preparation W (0.34 g, 2.5 mmol) and pyridine (0.60 g, 7.6 mmol) in CH₂Cl₂ (50 mL). The resulting suspension was stirred 20 h, filtered, and extracted with 5% Na₂CO₃ (30 mL). The organic layer was dried (Na₂SO₄) and concentrated to dryness to give a brown solid which was purified by preparative TLC R_f 0.58 (Whatman PLK5F, 10% CH₃OH/CH₂Cl₂) to give the title compound (0.38 g, 74%): MS (ES) m/e 413 [M+H]⁺.

b) (±)-7-[[[(2-Amino-4-pyrimidinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid

A solution of the compound of Example 4(a) in a mixture of CH₃OH (30 mL) and 0.5 N NaOH (6.0 mL) was heated at 50°C for 2 h, cooled to RT, and treated with TFA (1.0 mL). The solution was evaporated to dryness and the residue was purified by HPLC t_R 21.0 min (ODS-AQ, 50 X 250 mm, 90 mL/min, 86:14 CH₃CN:H₂O-0.1% TFA, UV detection at 220 nm) to give the title compound (0.150 g): MS (ES) m/e 399[M+H]⁺.

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Example 5Preparation of 3-[3,4-Dihydro-8-[[[(6-amino-2-pyridinyl)methyl]methylamino]carbonyl]-1-methyl-2,5-dioxo-1H-1,4-benzodiazepine]-4-propanoic acid

5

a) Benzyl 3-[3,4-dihydro-8-[[[(6-amino-2-pyridinyl)methyl]methylamino]carbonyl]-1-methyl-2,5-dioxo-1H-1,4-benzodiazepine]-4-propanoate

EDC (0.25 g, 1.3 mmol) is added to a solution of the compound of Preparation A(f) (1.1 mmol), the compound of Preparation T (1.1 mmol), HOBT·H₂O (170 mg, 1.3 mmol), and DIEA (0.9 mL, 4.4 mmol) in anhydrous CH₃CN (5 mL) at RT. After 21 h, the reaction is concentrated and the residue is purified by chromatography (silica gel) to afford the title compound.

10

b) 3-[3,4-Dihydro-8-[[[(6-amino-2-pyridinyl)methyl]methylamino]carbonyl]-1-methyl-2,5-dioxo-1H-1,4-benzodiazepine]-4-propanoic acid

15

A mixture of the compound of Example 5(a) (2 mmol) and 10% Pd/C (0.02 g) in EtOH (100 mL) is hydrogenated in an atmosphere of H₂ (50 psi) for 6 h. The catalyst is removed by filtration, and the filtrate is concentrated to afford the title compound.

Example 6

20

Preparation of 3-[4H-imidazo[1,2-a][1,4]benzodiazepine-5(6H)-1-methyl-6-oxo-9-[[[(6-amino-2-pyridinyl)methyl]methylamino]carbonyl]-5-propanoic acid

a) Ethyl 3-[4H-imidazo[1,2-a][1,4]benzodiazepine-5(6H)-1-methyl-6-oxo-9-[[[(6-amino-2-pyridinyl)methyl]methylamino]carbonyl]-5-propanoic acid

25

EDC (0.25 g, 1.3 mmol) is added to a solution of the compound of Preparation B(e) (1.1 mmol), the compound of Preparation T (1.1 mmol), HOBT·H₂O (170 mg, 1.3 mmol), and DIEA (0.9 mL, 4.4 mmol) in anhydrous CH₃CN (5 mL) at RT. After 21 h, the reaction is concentrated and the residue is purified by chromatography (silica gel) to afford the title compound.

30

b) 3-[4H-imidazo[1,2-a][1,4]benzodiazepine-5(6H)-1-methyl-6-oxo-9-[[[(6-amino-2-pyridyl)methyl]-N-methylamino]carbonyl]-5-propanoic acid

- 1M LiOH (3.8 mL, 3.8 mmol) is added dropwise to a solution of the compound of
5 Example 6(a) (2.5 mmol) in 1:1 CH₃OH:THF (20 mL) at RT. The resulting mixture is stirred for 20 h and concentrated. The residue is dissolved in H₂O and acidified with TFA (20%) to afford the title compound after chromatography.

Example 7

10

Preparation of 4-[4-[2-(6-Amino-2-pyridinyl)ethyl]-1-piperazinyl]-1-piperidineacetic acid

- a) Ethyl 4-[4-[2-[6-(phthalimido)-2-pyridinyl]ethyl]-1-piperazinyl]-1-(piperidinyl)acetate
According to the methods of Shoeb, et. al., Pharmazie, **1978**, 33, 581, and
15 Finkelstein, et. al., J. Amer. Chem. Soc., **1951**, 73, 302, a mixture compounds of Preparation C(b), Preparation D, and formaldehyde in EtOH is heated to reflux for 1 h, cooled, diluted with aqueous NaHCO₃, and extracted with EtOAc. The combined organic extracts are dried, evaporated, and flash chromatographed to give the title compound

- 20 b) 4-[4-[2-[6-(Phthalimido)-2-pyridinyl]ethyl]-1-piperazinyl]-1-piperidineacetic acid
A solution of Example 7(a) in NaOH /CH₃OH is stirred at RT for 18 h. The mixture is neutralized with HOAc and filtered to give the title compound.

- c) 4-[4-[2-(6-Amino-2-pyridinyl)ethyl]-1-piperazinyl]-1-piperidineacetic acid
25 A solution of Example 7(b) and hydrazine hydrate in EtOH is heated to reflux for 2 h.. The mixture is neutralised, desalted, and lyophilized to give the title compound.

Example 8

- 30 Preparation of 1-Hydroxy-4-[4-[(6-amino-2-pyridinyl)methyl]-1-piperazinyl]-cyclohexaneacetic acid

a) 1,1-Dimethylethyl 1-hydroxyl-4-[4-[(6-phthalimido-2-pyridinyl)methyl]-1-piperazinyl]-cyclohexaneacetate

5 A mixture 1,1-dimethylethyl 1-hydroxyl-4-(1-piperazinyl)-cyclohexaneacetate, Porter, et. al., EP 0 537 980 A1, 6-bromomethyl-2-(phthalimido)pyridine, US 4,490,533, and NaHCO₃ in CH₃CN is warmed. The mixture is concentrated and the residue is partitioned between H₂O and EtOAc. The organic phase is dried (Na₂SO₄), concentrated, and the residue is chromatographed (silica gel) to give the title compound.

10 b) 1-Hydroxyl-4-[4-[(6-phthalimido-2-pyridinyl)methyl]-1-piperazinyl]-cyclohexaneacetic acid

A solution of Example 8(b) in 4M HCl/dioxane/CH₂Cl₂ is stirred at RT for 18 h. The mixture is evaporated and filtered to give the title compound.

15 c) 1-Hydroxyl-4-[4-[(6-amino-2-pyridinyl)methyl]-1-piperazinyl]-cyclohexaneacetic acid

A solution of the compound of Example 8(b) in EtOH is treated with hydrazine hydrate and warmed. The mixture is concentrated and chromatographed (silica gel) to give the title compound.

20

Example 9

Preparation of 1-Hydroxyl-4-[4-[2-(6-amino-2-pyridinyl)ethyl]-1-piperazinyl]-cyclohexaneacetic acid

25 Following the general procedure of Example 7, except substituting 1,1-dimethylethyl 1-hydroxyl-4-(1-piperazinyl)-cyclohexaneacetate for the compound of Preparation C(b), gives the title compound.

Example 10Preparation of 4-[4-[(6-Amino-2-pyridinyl)methyl]-1-piperazinyl]-1-piperidineacetic acid

Following the general procedure of Example 8, except substituting the compound
 5 of Preparation C(b) for 1,1-dimethylethyl 1-hydroxyl-4-(1-piperazinyl)-
 cyclohexaneacetate, gives the title compound.

Example 11

10 Preparation of N-[[1-[[2-(6-Amino-2-pyridinyl)ethyl]carbonyl]-3-piperidinyl]carbonyl]-
 β-alanine

Following the procedures of Beavers et. al., WO 95/25091, Example 1, except
 substituting (6-amino-2-pyridinyl)propionic acid, Bondinell, et. al., WO 94/14775, for
 N^α-Boc-D-lys(Cbz)-OH, gives the title compound.

15

Example 12

Preparation of 2-[[6-Amino-2-pyridinyl)methyl]-5-[2-(carboxy-ethyl)amino]carbonyl]-
 -2,3-dihydro-3-oxo-1H-isoindole

20 Following the procedures of Preparation 1-12 in Hartman, et. al., EP 0 540 334
 A1, for the preparation of 2,3-dihydro-N-(2-carboxy-ethyl)-2-[2-(piperidinyl)ethyl]-3-
 oxo-1H-isoindole-5-carboxamide, except substituting the compound of Preparation U for
 Boc-4-piperidine-2-ethylamine, gives the title compound.

25

Example 13

Preparation of (S)-2-(Butylsulfonylamino)-3-[4-[[3-(6-amino-2-
 pyridinyl)propyl]oxy]phenyl]propionic acid

Following the procedures of Egbertson, et al., EP 0478363 A2, for the preparation of
 30 2-S-(butylsulfonylamino)-3-[4-(N-benzyloxycarbonylpiperidin-4-yl)-2,2-
 dimethyl]butyloxyphenylpropionic acid, except substituting the compound of Preparation
 E(b) for 4-[4-(N-benzyloxycarbonylpiperidin-4-yl)-2-methyl]pentan-2-ol in , gives the
 title compound.

Example 14Preparation of N-[3(R)-[2-(6-Amino-2-pyridinyl)ethyl]-2-oxopiperidinyl]acetyl]-3(R)-methyl-β-alanine

- 5 Following the procedure of Duggan, et. al., J. Med. Chem. 1995, 38, 3332, except substituting (6-amino-2-pyridinyl)butanoic acid, Bondinell, et. al., WO 94/14775, for (N-Boc-piperidin-4-yl)butanoic acid, gives the title compound.

Example 15

10

Preparation of 3-[[[3-[2-(6-Aminopyrid-2-yl)ethyl]isoxazolin-5(R,S)-yl]acetyl]aminol-3(R,S)-methylpropanoic acid

- 15 a) 4-[6-(Toluenesulfonylamino)-2-pyridinyl]-1-butanoximinoyl chloride
Following the procedure of Example 1(b) in WO 95/14682, except substituting the compound of Preparation F(e) for the 4-cyanobenzoxime, the title compound is prepared.

- 20 b) tert-Butyl [3-[2-[(6-toluenesulfonylamino)-2-pyridinyl]ethyl]isoxazolin-5(R,S)-yl]acetate

Following the procedure of Example 1(d) in WO 95/14682, except substituting the compound of Example 15(a) for 4-cyanobenzoximinoyl chloride, and substituting tert-butyl 3-butenate for methyl 3-butenate, the title compound is prepared.

- 25 c) [3-[2-[(6-Toluenesulfonylamino)-2-pyridinyl]ethyl]isoxazolin-5(R,S)-yl]acetic acid
4 M HCl in dioxane (10 mL) is added to a solution of the compound of Example 15(b) (5 mmol) in CH₂CH₂ (40 mL) at 0°C. The reaction is stirred at RT until complete, then is concentrated to afford the title compound.

d) Ethyl 3-[[3-[2-(6-aminopyrid-2-yl)ethyl]isoxazolin-5(R,S)-yl]acetyl]amino-3(R,S)-methylpropanoate

EDC (1.2 mmol) is added to a solution of the compound of Example 15(c) (1 mmol), ethyl 3(R,S)-aminobutyrate (1.2 mmol), HOBt · H₂O (1.2 mmol), and DIEA (4 mmol) in anhydrous CH₃CN (5 mL) at RT. The reaction is stirred at RT until complete, then is concentrated, and the residue is purified by chromatography (silica gel) to afford the title compound.

e) 3-[[[3-[2-(6-Aminopyrid-2-yl)ethyl]isoxazolin-5(R,S)-yl]acetyl]amino]-3(R,S)-methylpropanoic acid

1.0 N LiOH (2.5 mmol) is added to a solution of the compound of Example 15(d) (0.5 mmol) in THF (2.5 mL). The reaction is stirred at RT until complete, then is neutralized with 1.0 N HCl. The solution is concentrated and the residue is purified by reverse-phase chromatography to afford the title compound.

Example 16

Preparation of N-[3-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]benzoyl]-β-alanine

a) Benzyl N-[3-[[[(6-amino-2-pyridinyl)methyl]carbonyl]amino]benzoyl]-β-alaninate

A mixture of benzyl N-(3-aminobenzoyl)-β-alaninate, Alig, et. al., EP 0372486, (1 mmol), the compound of Preparation G (1 mmol), EDC (1.5 mmol), and DIEA (3 mmol) in DMF (25 mL) is stirred at RT. The mixture is poured into 5% NaHCO₃ and extracted with EtOAc. The combined organic phase is washed with H₂O, dried (MgSO₄), and concentrated. The residue is chromatographed (silica gel) to give the title compound.

b) N-[3-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]benzoyl]-β-alanine

A mixture of the compound of Example 16(b) (1 mmol) and 1N NaOH (1.5 mL) in CH₃OH (20 mL) is stirred and concentrated. The residue is dissolved in H₂O, extracted with CH₂Cl₂, and the aqueous phase is adjusted to pH 5 with dilute HCl to give the title compound.

Example 17

5 Preparation of [[1-[N-[(6-Amino-2-pyridinyl)methyl]carbonyl]tyrosyl]-4-piperidinyl]oxy]acetic acid

a) tert-Butyl [[1-[N-[(6-amino-2-pyridinyl)methyl]carbonyl]tyrosyl]-4-piperidinyl]oxy]acetate

10 A mixture of tert-butyl [(1-tyrosyl-4-piperidinyl)oxy]acetate, Alig, et. al., EP 372486, (1 mmol), the compound of Preparation G (1 mmol), EDC (1.5 mmol), and DIEA (3 mmol) in DMF (25 mL) is stirred at RT. The mixture is poured into 5% NaHCO₃ and extracted with EtOAc. The combined organic phase is washed with H₂O, dried (MgSO₄), and concentrated. The residue is chromatographed (silica gel) to give the title compound.

15

b) [[1-[N-[(6-Amino-2-pyridinyl)methyl]carbonyl]tyrosyl]-4-piperidinyl]oxy]acetic acid

A mixture of the compound of Example 17(a) (1 mmol) and CF₃CO₂H in CH₂Cl₂ is stirred and concentrated to give the title compound.

20

Example 18

25 Preparation of (±)-3-[[[2-(6-Aminopyrid-2-yl)ethyl]amino]succinoyl]amino]-4-pentynoic acid

a) Methyl 4-[[2-(6-aminopyrid-2-yl)ethyl]amino]-4-oxobutyrates

30 3-Carbomethoxypropionyl chloride (0.74 mL, 6.0 mmol) is added at 0°C to a stirred solution of the compound of Preparation H (5.0 mmol) and DIEA (4.4 mL, 25 mmol) in dry CH₂Cl₂ (50 mL). After stirring for 1.5 hr at RT, the reaction mixture is diluted with CH₂Cl₂ (50 mL) and washed sequentially with H₂O (25 mL) and 5%

NaHCO₃ (25 mL). The organic layer is dried (MgSO₄), concentrated, and reconcentrated from toluene. Chromatography (silica gel) gives the title compound.

b) 4-[[2-(6-Aminopyrid-2-yl)ethyl]amino]-4-oxobutyric acid

5 A mixture of the compound of Example 18(a) (530.6 mg, 2.0 mmol), 1.0 N LiOH (3.0 mL, 3.0 mmol), THF (10 mL), and H₂O (7 mL) is stirred at RT overnight, then is concentrated to dryness. The residue is taken up in H₂O (5 mL) and neutralized with 1.0 N HCl. The precipitate is collected and dried in vacuum to give the title compound.

10 c) Ethyl (±)-3-[[[2-(6-aminopyrid-2-yl)ethyl]amino]succinoyl]amino]-4-pentynoate

EDC (230 mg, 1.2 mmol) is added to a solution of the compound of Example 18(b) (203.3 mg, 1.0 mmol), ethyl (±)-3-amino-4-pentynoate, WO 93/07867, (169.4 mg, 1.2 mmol), HOBT · H₂O (162.2 mg, 1.2 mmol), and DIEA (0.70 mL, 4 mmol) in anhydrous CH₃CN (5 mL) at RT. The reaction is stirred at RT overnight, then is
15 concentrated to dryness. The residue is chromatographed (silica gel) to give the title compound.

d) (±)-3-[[[2-(6-Aminopyrid-2-yl)ethyl]amino]succinoyl]amino]-4-pentynoic acid

A mixture of the compound of Example 18(c) (187.2 mg, 0.5 mmol), 1.0 N LiOH
20 (0.75 mL, 0.75 mmol), THF (2.5 mL), and H₂O (1.7 mL) is stirred at RT overnight, then is concentrated to dryness. The residue is taken up in H₂O (2 mL) and acidified with TFA. ODS chromatography followed by lyophilization of the purified material gives the title compound.

25

Example 19

Preparation of (S)-4-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]glycyl]-3-methoxycarbonylmethyl-2-oxopiperazine-1-acetic acid

Following the procedure of Sugihara, et. al., EP 0529858, Example 59, except
30 substituting the compound of Preparation G for 4-amidinobenzoic acid hydrochloride, gives the title compound.

Example 20

Preparation of (3S,5S)-5-[4-[(6-Amino-2-pyridinyl)methyl]phenyl]oxymethyl]-3-carboxymethyl-2-pyrrolidinone

a) (3S,5S)-5-[4-[(6-Amino-2-pyridinyl)methyl]phenyl]oxymethyl]-3-[(tert-butoxycarbonyl)methyl]-2-pyrrolidinone

Following the procedure of Himmelsbach, et.al., AU-A-86926/91, Example 3(51), except substituting the compound of Preparation I for 4'-cyano-3'-fluoro-4-(hydroxy)biphenyl, gives the title compound.

b) (3S,5S)-5-[4-[(6-Amino-2-pyridinyl)methyl]phenyl]oxymethyl]-3-carboxymethyl-2-pyrrolidinone

Following the procedure of Himmelsbach, et. al., AU-A-86926/91, Example 7(93), except substituting the compound of Example 20(a) for (3S,5S)-3-[(tert-butyloxycarbonyl)methyl]-5-[(4'-amidino-3'-fluoro-4-biphenyl)oxymethyl]-2-pyrrolidinone, gives the title compound.

Example 21

Preparation of 1-[(6-Amino-2-pyridinyl)methyl]-3-[4-(2-carboxyethyl)phenyl]-4-methoxy-3-pyrrolin-2-one

Following the procedures of Linz, et. al., EP 0567968, except substituting the compound of Preparation U for 4-cyanoaniline, gives the title compound.

Example 22

Preparation of 4-[[[(6-Amino-2-pyridinyl)methyl]methylamino]acetyl]phenoxyacetic acid

a) Methyl 4-[[[(6-amino-2-pyridinyl)methyl]methylamino]acetyl]phenoxyacetate

Following the procedure of Wayne et. al., WO 94/22834, Example 1, except substituting the compound of Preparation T (1 mmol) for 1-(4-pyridyl)piperazine, gives the title compound.

5

b) 4-[[[(6-Amino-2-pyridinyl)methyl]methylamino]acetyl]phenoxyacetic acid

Following the procedure of Wayne et. al., WO 94/22834, Example 2, except substituting the compound of Example 22(a) for methyl 4-[2-[4-(4-pyridinyl)piperazin-1-yl]acetyl]phenoxyacetate, gives the title compound.

10

Example 23

Preparation of 2,2'-[[4-[[[(6-Amino-2-pyridinyl)methyl]methylamino]acetyl]-1,2-phenylene]bis(oxy)]bis-acetic acid

15

a) Dimethyl 2,2'-[[4-[[[(6-amino-2-pyridinyl)methyl]methylamino]acetyl]-1,2-phenylene]bis(oxy)]bis-acetate

Following the procedure of Wayne et. al., WO 94/22834, Example 3, except substituting the compound of Preparation T for 1-(4-pyridyl)piperazine, gives the title compound.

20

b) 2,2'-[[4-[[[(6-Amino-2-pyridinyl)methyl]methylamino]acetyl]-1,2-phenylene]bis(oxy)]bis-acetic acid

Following the procedure of Wayne et. al., WO 94/22834, Example 4, except substituting the compound of Example 23(a) for dimethyl 2,2'-[4-[2-[4-(4-pyridinyl)piperazin-1-yl]acetyl]phenylene-1,2-dioxy]diacetate, gives the title compound.

25

Example 24

30 Preparation of 4-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]methylamino]acetyl]phenoxyacetate

a) Benzyl 4-[[[(6-amino-2-pyridinyl)methyl]carbonyl] methylamino] acetyl] phenoxyacetate

A mixture of the compound of Preparation J(c) (1 mmol), Preparation G (1 mmol), EDC (1.5 mmol), and DIEA (3 mmol) in DMF (25 mL) is stirred at RT. The mixture is poured in to 5% NaHCO₃ and extracted with EtOAc. The organic phase is washed with H₂O, dried (MgSO₄), and concentrated. The residue is chromatographed (silica gel) to give the title compound.

b) 4-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]methylamino]acetyl]phenoxyacetate

The compound of Example 24(a) (1 mmol) and 1N NaOH (1.5 mL) in CH₃OH (20 mL) is stirred and concentrated. The residue is dissolved in H₂O, extracted with CH₂Cl₂, and the aqueous phase is adjusted to pH 5 with dilute HCl to give the title compound.

Example 25

Preparation of 4-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]methylamino]acetyl]-1,2-phenylenedioxydiacetic acid

a) Dimethyl 4-[[[(6-amino-2-pyridinyl)methyl]carbonyl]methylamino]acetyl]-1,2-phenylenedioxydiacetate

Following the procedure of Example 24(a), except substituting the compound of Preparation K(c) for the compound of Preparation J(c), gives the title compound.

b) 4-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]methylamino]acetyl]-1,2-phenylenedioxydiacetic acid

Following the procedure of procedure of Example 24(b), except substituting the compound of Example 25(a) for the compound of Example 24(a), gives the title compound.

Example 26Preparation of 1-[(2-Amino-6-pyridinyl)methyl]-3-[4[2-(carboxy)ethyl]phenyl]-3-oxo-imidazolidine

5

a) Ethyl 2-[4-(2-hydroxyethylamino)phenyl]propionate

Following the procedure of Himmelsbach, et. al., EP 0587134, Example V, glycolaldehyde dimer (Aldrich) (1 mmol) is added to a solution of methyl 2-(4-aminophenyl)propionate, (1 mmol) in aqueous CH₃CN (pH 6-7) (10 mL), followed by 10 NaBH₃CN (1.2 mmol), and the mixture is allowed to stir for 1 h. The mixture is concentrated to an oil, and the residue is dissolved in a mixture of ice water and EtOAc. The aqueous layer is neutralized with 4 N NaOH and washed with EtOAc. The organic phase is concentrated and a solution of the resulting oil in EtOAc is chromatographed (silica gel, gradient, 5-30% CH₃OH/CH₂Cl₂-0.1% HOAc). The fractions containing the 15 product are combined and concentrated to give the title compound.

b) N-[(2-Phthaloyl-6-pyridinyl)methyl]-N'-hydroxyethyl-N'-[4[(2-ethoxycarbonyl)ethyl]]phenyl]-urea

Following the procedures of Himmelsbach, et. al., EP 0587134 and EP 0612741, a 20 solution of the compound of Preparation L (1 mmol) and COCl₂ (1.1 mmol) in THF (20 mL) is allowed to stir at -20°C for 20 min. The compound of Example 26(a) (1 mmol) is added to the solution and the resulting mixture is allowed to stir and warm RT for 18 h. The resulting solution is concentrated and a solution of the resulting residue in EtOAc is washed with 5% citric acid followed by H₂O. The organic phase is concentrated and a 25 solution of the resulting oil in EtOAc is chromatographed (silica gel, gradient, 5-30% CH₃OH/CH₂Cl₂-0.1% HOAc). The fractions containing the product are combined and concentrated to give the title compound.

c) N¹-[(2-Phthaloyl-6-pyridinyl)methyl]-N³-[4[(2-ethoxycarbonyl)ethyl]]phenyl]-2-oxo-imidazolidine

Following the procedures of Himmelsbach, et. al., EP 0587134, Example III, and EP 0612741, a solution of the compound of Example 26(b) (1 mmol), methanesulfonyl chloride (1.2 mmol) and Et₃N (1.2 mmol) in CH₂Cl₂ (5 mL) is allowed to stir at 0°C for 1 h. The mixture is partitioned between H₂O and CH₂Cl₂. The organic phases are combined, dried (Na₂SO₄), and concentrated. A solution of the residue and NaI (1.1 mmol) in acetone (5 mL) is heated to reflux for 3 h and then concentrated. Potassium bis(trimethylsilyl)azide (1.2 mmol) is added to a solution of the resulting oil in DMF (5 mL), cooled to 0°C. The solution is allowed to warm to RT over 30 min and concentrated to give an oil. The oil is partitioned between H₂O and CH₂Cl₂. The organic phases are combined, dried (Na₂SO₄), and concentrated. A solution of the resulting oil in EtOAc is chromatographed (silica gel, gradient, 5-30% CH₃OH/CH₂Cl₂-0.1% HOAc). The fractions containing the product are combined and concentrated to give the title compound.

d) N¹-[(Amino-6-pyridinyl)methyl]-N³-[4[(2-carboxyl)ethyl]]phenyl]-2-oxo-imidazolidine

Following the procedures of Himmelsbach, et. al., EP 0587134, Example III, and EP 0612741, a solution of the compound of Example 26(c) (1 mmol) and hydrazine hydrate (1.1 mmol) in EtOH (10 mL) is allowed to stir for 18 h. The solution is concentrated and the residue is partitioned between H₂O and EtOAc. The organic phases are combined and concentrated. A solution of the resulting oil in THF (5 mL) and 1 N NaOH (1.2 mL, 1.2 mmol) is allowed to stir for 18 h. The mixture is neutralized with conc HCl and chromatographed (silica gel, gradient, 5-30% CH₃OH/CH₂Cl₂-0.1% HOAc). The fractions containing the product are combined and concentrated to give the title compound.

Example 27Preparation of [6-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-1,2,3,4-tetrahydroisoquinolin-2-yl]acetic acid

5

a) Ethyl [6-[[[(6-amino-2-pyridinyl)methyl]methylamino]carbonyl]-1,2,3,4-tetrahydroisoquinolin-2-yl]acetate

A solution of the compound of Preparation M(d) (0.263 g, 1.0 mmol), the compound of Preparation T (0.34 g, 1.0 mmol), EDC (0.191 g, 1.0 mmol), HOBt (0.151 g, 1.0 mmol) and Et₃N (0.235 mL, 2.0 mmol) in DMF (7 mL) is stirred for 8 h. The solution is concentrated to an oil which is purified by chromatography (silica gel, gradient, 10-33% CH₃OH/CH₂Cl₂) to afford the title compound (0.32 g, 77%)

10

b) [6-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-1,2,3,4-tetrahydroisoquinolin-2-yl]acetic acid

15

A solution of the compound of Example 27(a) (0.42 g, 1.0 mmol) in 1N NaOH (1.5 mL, 1.5 mmol) and EtOH (5 mL) is stirred for 8 h. The solution is concentrated to an oil which is purified by chromatography (silica gel, gradient, 10-33% CH₃OH/CH₂Cl₂) to afford the title compound (0.35 g, 76%).

20

Example 28Preparation of [6-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-1,2,3,4-tetrahydro-1-oxo-isoquinolin-2-yl]acetic acid

25

a) Ethyl [6-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-1,2,3,4-tetrahydro-1-oxo-isoquinolin-2-yl]acetate

Following the procedure of Example 27(a), except substituting the compound of Preparation N(d) for the compound of Preparation M(d), gives the title compound.

30

Alternatively, a solution of the compound of Preparation N(c) (0.23 g, 1.0 mmol), Pd(OAc)₂ (0.026 g, 0.1 mmol), the compound of Preparation T (0.31 g, 1.0 mmol), Ph₃P (0.262 g, 1.0 mmol), diisopropylamine (0.25 mL, 2.1 mmol), and NMP (7 mL) in 10% NH₄CO₃ is stirred for 8 h under an atmosphere of CO. The solution is concentrated to an oil which is purified by chromatography (silica gel, gradient, 25-75% CH₃OH/CH₂Cl₂) to afford the title compound (0.26 g, 76%)

b) [6-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-1,2,3,4-tetrahydro-1-oxo-isoquinolin-2-yl]acetic acid

Following the procedure of Example 27(b), except substituting the compound of Example 28(a) for the compound of Example 27(a), gives the title compound.

Example 29

Preparation of [6-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]tetralin-2-yl]acetic acid

a) tert-Butyl [6-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]tetralin-2-yl]acetate

Following the procedure of Example 27(a), except substituting tert-butyl (6-amino-tetralin-2-yl)acetate [Fisher, et. al., EO 0635492, Scheme 12 and Example 28, parts A-D] for the compound of Preparation M(d) and substituting the compound of Preparation G for the compound of Preparation T, gives the title compound.

b) [6-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]tetralin-2-yl]acetic acid

A solution of the compound of Example 29(a) (0.32 g, 1.0 mmol) and TFA (5 mL) in CH₂Cl₂ (5 mL) is allowed to stir for 1 h. The solution is concentrated to an oil which is treated with Et₂O. Filtration and drying *in vacuo* afforded the title compound (0.15g, 50%)

Example 30Preparation of [6-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]tetralin-2-yl]acetic acid

- 5 Following the procedure of Example 27, except substituting the compounds of Preparation O(b) for the compound of Preparation M(d), gives the title compound.

Example 31

10 Preparation of [5-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]benzofuran-2-yl]propionic acid

 Following the procedure of Example 27, except substituting ethyl (5-aminobenzofuran-2-yl)propionate from Preparation P(e) for the compound of Preparation M(d), gives the title compound.

15

Example 32

Preparation of [5-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]-2,3-dihydro-benzofuran-2-yl]propionic acid

- 20 Following the procedure of Example 27, except substituting ethyl (5-amino-2,3-dihydro-benzofuran-2-yl)propionate from Preparation P(e) for the compound of Preparation M(d), gives the title compound.

Example 33

25

Preparation of [5-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]benzofuran-2-yl]propionic acid

- Following the procedure of Example 28, except substituting the compounds of Preparation Q(d) or (e) for the compounds of Preparation N(c) or (d), gives the title
- 30 compound.

Example 34Preparation of [5-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-2,3-dihydro-benzofuran-2-yl]-propionic acid

- 5 Following the procedure of Example 28, except substituting the compounds of Preparation R(a) or (b) for the compounds of Preparation N(c) or (d), gives the title compound.

Example 35

10 Preparation of (±)-3-[[[4-(6-Amino-2-pyridinyl)butanoyl]glycyl]amino]-4-pentynoic acid

- a) Ethyl (±)-3-[[[4-(6-Amino-2-pyridinyl)butanoyl]glycyl]amino]-4-pentynoate

 DIEA (5.43 mmol) is added to a stirred solution of the compound of Preparation S(b) (1.76 mmol), 4-(6-amino-2-pyridinyl)butyric acid, Bondinell, et. al., WO 94/14775, (1.55 mmol), HOBT · H₂O (2.33 mmol), and EDC (2.33 mmol) in anhydrous CH₃CN (15 mL) at RT. The reaction mixture is stirred, concentrated, diluted with CH₂Cl₂ (100 mL), and washed sequentially with 5% NaHCO₃ and brine. Drying (MgSO₄), concentration, and chromatography (silica gel, CH₃OH/ CH₂Cl₂) gives the title compound.

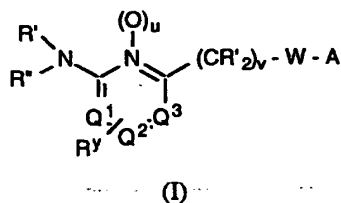
- 20 b) (±)-3-[[[4-(6-Amino-2-pyridinyl)butanoyl]glycyl]amino]-4-pentynoic acid

 1.0 N LiOH (0.71 mmol) is added dropwise at RT to a mixture of the compound of Example 35(a) (0.285 mmol) in THF (5 mL), H₂O (5 mL) and CH₃CN (1 mL). The mixture is stirred, concentrated to a small volume, and cooled in an ice bath before neutralizing with 1.0 N AcOH (0.70 mL). The solution is lyophilized and the residue is purified by chromatography (ODS, CH₃CN/H₂O-0.1% TFA) to give the title compound.

 The above description fully discloses how to make and use the present invention. However, the present invention is not limited to the particular embodiments described hereinabove, but includes all modifications thereof within the scope of the following claims. The various references to journals, patents and other publications which are cited herein comprises the state of the art and are incorporated herein by reference as though fully set forth.

What is claimed is:

1. A compound according to formula (I):



wherein

A is a fibrinogen antagonist template;

W is a linking moiety of the form $-(\text{CHR}^g)_a-\text{U}-(\text{CHR}^g)_b-\text{V}-$;

10 Q^1 , Q^2 and Q^3 are independently N or $\text{C}-\text{R}^y$, provided that no more than one of Q^1 , Q^2 and Q^3 is N;

R' is H or C_{1-6} alkyl, C_{3-7} cycloalkyl- C_{0-6} alkyl or Ar- C_{0-6} alkyl

R'' is R' , $-\text{C}(\text{O})\text{R}'$ or $-\text{C}(\text{O})\text{OR}'$;

R^g is H or C_{1-6} alkyl, Het- C_{0-6} alkyl, C_{3-7} cycloalkyl- C_{0-6} alkyl or Ar- C_{0-6} alkyl;

15 R^k is R^g , $-\text{C}(\text{O})\text{R}^g$ or $-\text{C}(\text{O})\text{OR}^g$

R^i is H, C_{1-6} alkyl, Het- C_{0-6} alkyl, C_{3-7} cycloalkyl- C_{0-6} alkyl, Ar- C_{0-6} alkyl, Het- C_{0-6} alkyl- U' - C_{1-6} alkyl, C_{3-7} cycloalkyl- C_{0-6} alkyl- U' - C_{1-6} alkyl or Ar- C_{0-6} alkyl- U' - C_{1-6} alkyl;

R^y is H, halo, $-\text{OR}^g$, $-\text{SR}^g$, $-\text{CN}$, $-\text{NR}^g\text{R}^k$, $-\text{NO}_2$, $-\text{CF}_3$, $\text{CF}_3\text{S}(\text{O})_r$, $-\text{CO}_2\text{R}^g$, $-\text{COR}^g$ or $-\text{CONR}^g_2$;

20 U and V are absent or CO, CR^g_2 , $\text{C}(\text{=CR}^g_2)$, $\text{S}(\text{O})_c$, O, NR^g , CR^gOR^g , $\text{CR}^g(\text{OR}^k)\text{CR}^g_2$, $\text{CR}^g_2\text{CR}^g(\text{OR}^k)$, $\text{C}(\text{O})\text{CR}^g_2$, $\text{CR}^g_2\text{C}(\text{O})$, CONR^i , NR^iCO , $\text{OC}(\text{O})$, $\text{C}(\text{O})\text{O}$, $\text{C}(\text{S})\text{O}$, $\text{OC}(\text{S})$, $\text{C}(\text{S})\text{NR}^g$, $\text{NR}^g\text{C}(\text{S})$, $\text{S}(\text{O})_2\text{NR}^g$, $\text{NR}^g\text{S}(\text{O})_2$, $\text{N}=\text{N}$, NR^gNR^g , NR^gCR^g_2 , NR^gCR^g_2 , CR^g_2O , OCR^g_2 , $\text{CR}^g=\text{CR}^g$, $\text{C}\equiv\text{C}$, Ar or Het;

25 a is 0, 1 or 2;

b is 0, 1 or 2;

c is 0, 1 or 2;

r is 0, 1 or 2;

u is 0 or 1; and

30 v is 0 or 1;

or pharmaceutically acceptable salts thereof;

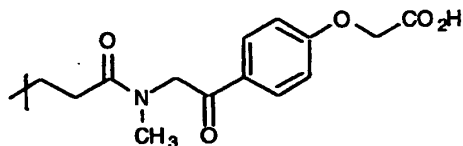
provided that:

(i) when v is 0, and R' , R'' and R' are H, and Q^1 - Q^3 are CH,

W-A is not 7-aminocarbonyl-2,3,4,5-tetrahydro-3-oxo-4-methyl-1H-1,4-benzodiazepine-2-acetic acid, 7-aminocarbonyl-1-acetyl-2,3,4,5-tetrahydro-3-oxo-4-methyl-1H-1,4-benzodiazepine-2-acetic acid, or 7-aminocarbonyl-2,3,4,5-tetrahydro-3-oxo-4-methyl-1H-1-benzazepine-2-acetic acid, or the methyl esters thereof;

(ii) when v is 0 or 1 and R' , R'' and R' are H, and Q^1 - Q^3 are CH,

W-A is not 3-propanoyl-glycyl-aspartyl-phenylalanine, or



, and the benzyl esters thereof.

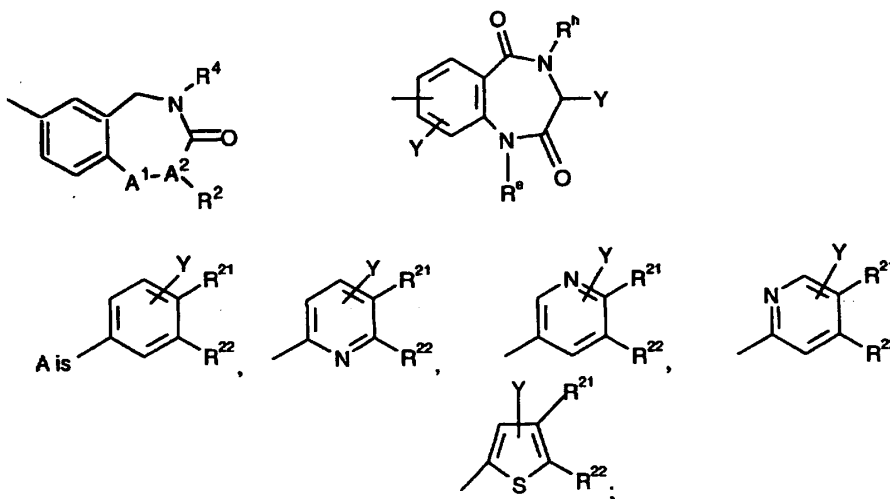
2. A compound according to claim 1 in which Q^1 , Q^2 and Q^3 are each CH, and u is 0.

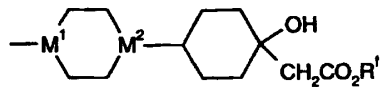
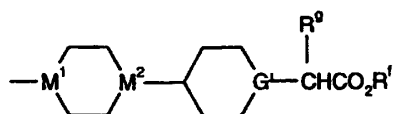
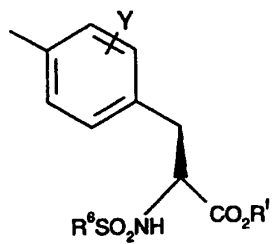
3. A compound according to claim 1 in which R' is H and R'' is H or C_{1-4} alkyl.

4. A compound according to claim 1 in which W is $-(CHR^8)_a-CONR^i-$ or

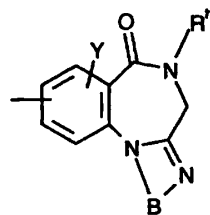
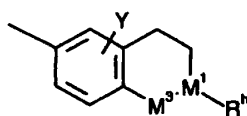
$-(CHR^8)_a-NR^iCO-$.

5. A compound according to formula (I) in which A is chosen from the group of

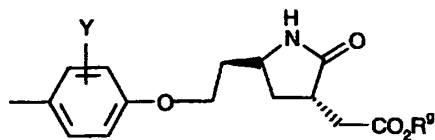
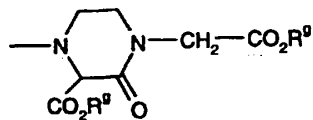
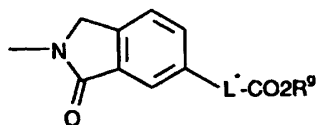




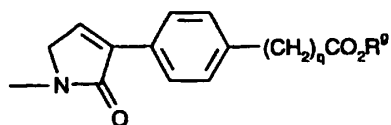
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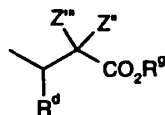
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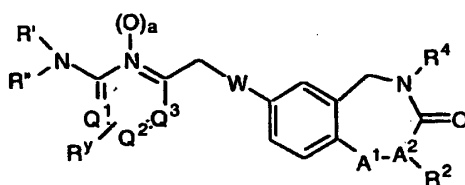


and



5 and which has 13-14 covalent bonds between the acidic moiety and the first nitrogen in the pyridine ring.

6. A compound according to claim 1 in which is:



10

wherein

A¹-A² is NR¹-CH, NC(O)R³-CH, N=C, CR¹=C, CHR¹-CH, O-CH or S-CH;

R¹ is H, C₁₋₆ alkyl or benzyl;

15 R² is (CH₂)_qCO₂H;

R⁴ is H, C₁₋₆alkyl, Ar-C₀₋₆alkyl, Het-C₀₋₆alkyl, or C₃₋₆cycloalkyl-C₀₋₆alkyl; and

q is 1, 2 or 3.

7. A compound according to claim 6 wherein A¹-A² is NH-CH and R² is CH₂CO₂H.

20

8. A compound according to claim 7 wherein W is -(CHR⁸)_a-CONRⁱ- or -(CHR⁸)_a-NRⁱCO-.

9. A compound according to claim 1 which is:

25 3-[3,4-Dihydro-8-[[[(6-amino-2-pyridinyl)methyl]methylamino]carbonyl]-1-methyl-2,5-dioxo-1H-1,4-benzodiazepine]-4-propanoic acid;

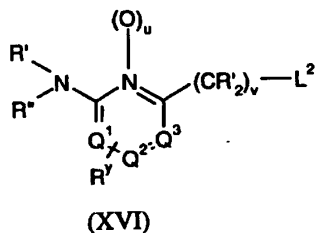
3-[4H-imidazo[1,2-a][1,4]benzodiazepine-5(6H)-1-methyl-6-oxo-9-[[[(6-amino-2-pyridinyl)methyl]methylamino]carbonyl]-5-propanoic acid;

4-[4-[2-(6-Amino-2-pyridinyl)ethyl]-1-piperazinyl]-1-piperidineacetic acid ;

- 1-Hydroxyl-4-[4-[(6-amino-2-pyridinyl)methyl]-1-piperazinyl]-cyclohexaneacetic acid;
 1-Hydroxyl-4-[4-[2-(6-amino-2-pyridinyl)ethyl]-1-piperazinyl]-cyclohexaneacetic acid;
 4-[4-[(6-Amino-2-pyridinyl)methyl]-1-piperazinyl]-1-piperidineacetic acid ;
 N-[[1-[[2-(6-Amino-2-pyridinyl)ethyl]carbonyl]-3-piperidinyl]carbonyl]-b-alanine;
 5 2-[(6-Amino-2-pyridinyl)methyl]-5-[2-(carboxy-ethyl)amino]carbonyl];
 -2,3-dihydro-3-oxo-1H-isoindole;
 (S)-2-(Butylsulfonylamino)-3-[4-[[3-(6-amino-2-pyridinyl)propyl]oxy]phenyl]propionic
 acid;
 N-[3(R)-[2-(6-Amino-2-pyridinyl) ethyl]-2-oxopiperidinyl]acetyl]-3(R)-methyl-b-
 10 alanine;
 3-[[[3-[2-(6-Aminopyrid-2-yl)ethyl]isoxazolin-5(R,S)-yl]acetyl]amino]-3(R,S)-
 methylpropanoic acid;
 N-[3-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]benzoyl]-b-alanine;
 [[1-[N-[[[(6-Amino-2-pyridinyl))methyl]carbonyl]tyrosyl]-4-piperidinyl]oxy]acetic acid ;
 15 (±)-3-[[[2-(6-Aminopyrid-2-yl)ethyl]amino]succinoyl]amino]-4-pentynoic acid;
 (S)-4-[[[(6-Amino-2-pyridinyl)methyl]carbonyl]glycyl]-3-methoxycarbonylmethyl-2-
 oxopiperazine-1-acetic acid ;
 (3S,5S)-5-[4-[(6-Amino-2-pyridinyl)methyl]phenyl]oxymethyl]-3-carboxymethyl-2-
 pyrrolidinone;
 20 1-[(6-Amino-2-pyridinyl)methyl]-3-[4-(2-carboxyethyl)phenyl]-4-methoxy-3-pyrrolin-2-
 one;
 4-[[[(6-Amino-2-pyridinyl)methyl]methylamino]acetyl]phenoxyacetic acid;
 2,2'-[[4-[[[(6-Amino-2-pyridinyl)methyl]methylamino]acetyl]-1,2-
 phenylene]bis(oxy)]bis-acetic acid;
 25 4-[[[[[(6-Amino-2-pyridinyl)methyl]carbonyl]methylamino]acetyl]phenoxyacetate;
 4-[[[[[(6-Amino-2-pyridinyl)methyl]carbonyl]methylamino]acetyl]-1,2-
 phenylenedioxydiacetic acid;
 1-[(2-Amino-6-pyridinyl)methyl]-3-[4[2-(carboxy)ethyl]phenyl]-3-oxo-imidazolidine;
 [6-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-1,2,3,4-
 30 tetrahydroisoquinolin-2-yl]acetic acid;
 [6-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-1,2,3,4-tetrahydro-1-oxo-
 isoquinolin-2-yl]acetic acid;
 [6-[[[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]tetralin-2-yl]acetic acid;
 [6-[[[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]tetralin-2-yl]acetic acid;
 35 [5-[[[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]benzofuran-2-yl]propionic acid;
 [5-[[[[[(6-Amino-2-pyridinyl)methyl]carbonyl]amino]-2,3-dihydro-benzofuran-2-
 yl]propionic acid;

- [5-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]benzofuran-2-yl]propionic acid;
- [5-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-2,3-dihydro-benzofuran-2-yl]-propionic acid; or
- 5 (±)-3-[[[4-(6-Amino-2-pyridinyl)butanoyl]glycyl]amino]-4-pentynoic acid.
10. A compound according to claim 1 which is:
- (S)-7-[[[(6-Amino-2-pyridinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic;
- 10 (S)-7-[[[(6-Amino-2-pyridinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid;
- (S)-7-[[[(6-Ethylamino-2-pyridinyl)methyl]amino]carbonyl]-2,3,4,5-tetrahydro-4-methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid; or
- (±)-7-[[[(2-Amino-4-pyrimidinyl)methyl]methylamino]carbonyl]-2,3,4,5-tetrahydro-4-
- 15 methyl-3-oxo-1H-1,4-benzodiazepine-2-acetic acid.
11. A pharmaceutical composition which comprises a compound according to any one of claims 1-10 and a pharmaceutically acceptable carrier.
- 20 12. A method of treating a disease state in which antagonism of the vitronectin receptor is indicated which comprises administering a compound according to claim 1.
13. A method according to claim 12 for inhibiting angiogenesis or treating atherosclerosis, restenosis, inflammation, cancer or osteoporosis.
- 25 14. The use of a compound according to any one of claims 1-10 in the manufacture of a medicament.
15. The use of a compound according to any one of claims 1-10 in the manufacture of a medicament for the inhibition of the vitronectin receptor in a mammal in need thereof.
- 30 16. The use of a compound according to any one of claims 1-10 in the manufacture of a medicament for the treatment of atherosclerosis, restenosis, inflammation, cancer or osteoporosis.
- 35

17. A process for preparing a compound of formula (I) as defined in claim 1, which process comprises reacting a compound of formula (XVI) with a compound of formula (XVII):



wherein:

Q^1 , Q^2 , Q^3 , R^y , R' , R'' , u , v and A are as defined in formula (I), with any reactive functional groups protected; and

L^1 and L^2 are groups which react to form a covalent bond in the moiety W as defined in formula (I);

and thereafter removing any protecting groups, and optionally forming a pharmaceutically acceptable salt.

INTERNATIONAL SEARCH REPORT

national application No.
PCT/US96/20744

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : Please See Extra Sheet.
US CL : 514/221;540/513
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/221;540/513

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
CASONLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 3,627,754 A (NING ET AL) 14 December 1971, see entire document, especially column 1.	1-17, parts

<input type="checkbox"/> Further documents are listed in the continuation of Box C.	<input type="checkbox"/> See patent family annex.
<p>* Special categories of cited documents:</p> <p>*A* document defining the general state of the art which is not considered to be of particular relevance</p> <p>*B* earlier document published on or after the international filing date</p> <p>*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>*O* document referring to an oral disclosure, use, exhibition or other means</p> <p>*P* document published prior to the international filing date but later than the priority date claimed</p>	<p>*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>*Z* document member of the same patent family</p>

Date of the actual completion of the international search 06 MARCH 1997	Date of mailing of the international search report 27 MAR 1997
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230	Authorized officer <i>Robert W. Ramsuer</i> ROBERT W. RAMSUER Telephone No. (703) 308-1235

INTERNATIONAL SEARCH REPORT

I. national application No.
PCT/US96/20744

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☒ Claims Nos.: Claims 1-17, parts
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

Please See Extra Sheet.
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

I national application No.
PCT/US96/20744

A. CLASSIFICATION OF SUBJECT MATTER: IPC (6):

A61K 31/44; C07D 401/02

BOX I. OBSERVATIONS WHERE CLAIMS WERE FOUND UNSEARCHABLE

2. Where no meaningful search could be carried out, specifically:

The multitude of variables and their permutations and combinations (e.g. W,A,U,V,Q1,Q2,Q3,Ar,Het,a,b,c,the provisos,etc.) result in claimed subject matter that is so broad in scope that it is rendered virtually incomprehensible and thus no meaningful search can be given. Note also that the claimed subject matter lacks a significant structural element qualifying as the special technical feature that clearly defines a contribution over the art. The subject matter claimed contains a N-C-N-C group which does not define a contribution over the prior art. Therefore, the first discernable invention as found in Example 1,(the compound therein, the pharmaceutical composition therewith, the method of preparation thereof and the method of treating osteoporosis therewith) has been searched.